

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 77.2131 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-10
Perfect score: 50
Sequence: 1 TRLTRKDLK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_23Sep04:*

1: Geneseqp1980s:*

2: Geneseqp1990s:*

3: Geneseqp2000s:*

4: Geneseqp2001s:*

5: Geneseqp2002s:*

6: Geneseqp2003as:*

7: Geneseqp2003bs:*

8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------|--------------------|
| 1 | 50 | 100.0 | 10 | 2 | AAY30691 Apo-B100 |
| 2 | 46 | 92.0 | 10 | 2 | AAY30690 Apo-B100 |
| 3 | 42 | 84.0 | 11 | 2 | AAW57205 Apo B bin |
| 4 | 42 | 84.0 | 13 | 2 | AAW57207 Apo B 100 |
| 5 | 42 | 84.0 | 15 | 2 | AAW41261 Apolipop |
| 6 | 42 | 84.0 | 15 | 2 | AAW96892 ApoB-100 |
| 7 | 42 | 84.0 | 20 | 6 | ABJ37575 Heparin b |
| 8 | 42 | 84.0 | 22 | 2 | AAW57208 Apo B 100 |
| 9 | 42 | 84.0 | 22 | 2 | AAW57209 Apo B 100 |
| 10 | 42 | 84.0 | 34 | 5 | AAE14541 Human apo |
| 11 | 42 | 84.0 | 36 | 2 | AAW96876 Nucleic a |
| 12 | 42 | 84.0 | 37 | 2 | AAW64587 Human apo |
| 13 | 42 | 84.0 | 37 | 2 | AAW96845 Nucleic a |
| 14 | 42 | 84.0 | 34 | 4 | ABJ37687 Peptide # |
| 15 | 42 | 84.0 | 34 | 4 | ABJ37687 Peptide # |
| 16 | 42 | 84.0 | 37 | 2 | AAW57204 Human liv |
| 17 | 42 | 84.0 | 37 | 2 | AAW57204 Human apo |
| 18 | 42 | 84.0 | 37 | 2 | AAW57204 Human apo |
| 19 | 42 | 84.0 | 2463 | 8 | AAW57400 Human apo |
| 20 | 42 | 84.0 | 3923 | 2 | AAW57400 Human apo |
| 21 | 42 | 84.0 | 4536 | 2 | AAW57400 Human apo |
| 22 | 42 | 84.0 | 4536 | 2 | AAW57400 Human apo |
| 23 | 42 | 84.0 | 4560 | 5 | AAW57400 Human apo |
| 24 | 42 | 84.0 | 4561 | 7 | AAW57400 Human apo |
| 25 | 42 | 84.0 | 4563 | 5 | AAW57400 Human apo |

| | | | | | |
|----|------|------|------|---|----------|
| 26 | 42 | 84.0 | 4563 | 6 | ABU79140 |
| 27 | 42 | 84.0 | 4563 | 7 | ADP43408 |
| 28 | 42 | 84.0 | 4563 | 8 | ADH18871 |
| 29 | 42 | 84.0 | 4563 | 8 | ADH18870 |
| 30 | 42 | 84.0 | 4563 | 8 | ADH18870 |
| 31 | 42 | 84.0 | 4563 | 8 | ADH18870 |
| 32 | 42 | 84.0 | 4590 | 4 | AAU33184 |
| 33 | 39.5 | 79.0 | 11 | 2 | AAU33184 |
| 34 | 38 | 76.0 | 10 | 2 | AAU33184 |
| 35 | 38 | 76.0 | 10 | 2 | AAU33184 |
| 36 | 38 | 76.0 | 359 | 8 | ADL90227 |
| 37 | 38 | 76.0 | 390 | 7 | ADD44981 |
| 38 | 38 | 76.0 | 390 | 7 | ADD44985 |
| 39 | 38 | 76.0 | 390 | 7 | ADD44987 |
| 40 | 38 | 76.0 | 390 | 7 | ADD44983 |
| 41 | 38 | 76.0 | 390 | 7 | ADJ68663 |
| 42 | 38 | 76.0 | 390 | 8 | ADQ18720 |
| 43 | 38 | 76.0 | 397 | 5 | AAE27989 |
| 44 | 38 | 76.0 | 414 | 7 | ADJ68303 |
| 45 | 38 | 76.0 | 414 | 7 | ADJ68302 |

ALIGNMENTS

RESULT 1
AAY30691
ID AAY30691 standard; peptide; 10 AA.
XX
AC AAY30691;
XX
DT 17-NOV-1999 (first entry)
XX
DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS Synthetic.
OS Homo sapiens.
XX
FN WO9946598-A1.
XX
PD 16-SEP-1999.
XX
PF 05-MAR-1999; 99WO-US004805.
XX
PR 10-MAR-1998; 98US-0077618P.
XX (REGC) UNIV CALIFORNIA.
XX
PI Innerarity TL, Boren JOS;
XX
DR WPI; 1999-551509/46.
XX
PT Identifying compounds which affect binding of low density lipoprotein
PT with proteoglycan, used for, e.g. obtaining compounds for reducing
PT atherosclerosis.
XX
XX Claim 17; Page 57; 70pp; English.
XX
XX AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
XX receptor mutations. They were created to identify compounds which
XX modulate atherosclerosis. The peptides are derived from amino acids 3358
XX to 3367 of apoB100. The method comprises detecting compounds which affect
XX low density lipoprotein (LDL) binding with proteoglycan (PG). The method
XX can be used for identifying compounds which disrupt LDL-PG binding
XX without inhibiting LDL receptor binding. Such compounds can be used to
XX reduce or prevent the formation of atherosclerotic lesions and prevent
XX atherosclerosis. The transgenic non-human animals and mammals which
XX express human apo-B100 can be used as an in vivo model system for the
XX study of atherosclerosis, and in vivo assay methods for identifying
XX compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX

SQ Sequence 10 AA;

Query Match 100.0%; Score 50; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.003;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKDGLK 10
 |||||
 DB 1 TRLTRKDGLK 10

RESULT 2
 AAY30690
 ID AAY30690 standard; peptide; 10 AA.
 XX
 AC AAY30690;
 XX
 DT 17-NOV-1999 (first entry)
 XX
 DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
 XX
 KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN W09946598-A1.
 XX
 PD 16-SEP-1999.
 XX
 PF 05-MAR-1999; 99WO-US004805.
 XX
 PR 10-MAR-1998; 98US-0077618P.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 PI Innerarity TL, Boren JOS;
 XX
 PS WPI; 1999-551509/46.
 XX

XX Identifying compounds which affect binding of low density lipoprotein
 PT with proteoglycan, used for, e.g. obtaining compounds for reducing
 PT atherosclerosis.
 XX
 PS Claim 17; Page 57; 70pp; English.
 XX
 CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (Pg). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX

SQ Sequence 10 AA;
 Query Match 92.0%; Score 46; DB 2; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.02;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKDGLK 10
 |||||
 DB 1 TRLTRKEGLK 10

RESULT 3
 AAW57205
 ID AAW57205 standard; peptide; 11 AA.
 XX
 AC AAW57205;
 XX
 DT 03-AUG-1998 (first entry)
 XX
 DE Apo B binding site peptide 2.
 XX
 KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 OS Synthetic.
 XX
 PN W09813385-A2.
 XX
 PD 02-APR-1998.
 XX
 PF 25-SEP-1997; 97WO-GB002610.
 XX
 PR 27-SEP-1996; 96GB-00020153.
 XX
 PA (UYST) UNIV STRATHCLYDE.
 XX
 PI Halbert GW, Owens MD, Baillie G;
 XX
 DR WPI; 1998-230637/20.
 XX
 PT Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 PS Claim 12; Page 52; 73pp; English.
 XX

XX The present sequence represents a specifically claimed Apo B binding site
 CC peptide which can be used as a component of a non-naturally occurring,
 CC receptor-competent low density lipoprotein (LDL) particle of the present
 CC invention. The LDL particle comprises at least 1 peptide component that
 CC has at least 1 binding site for an apo B protein receptor and at least 1
 CC lipophilic substituent. Also described in the invention are peptides
 CC containing an apo B binding sequence with at least 70% identity with
 CC sequences: KASYKKNKHH (1) or TRLTRKRGK (2), or their dimers. Non-
 CC naturally occurring, receptor-competent LDL particles are useful as: (i)
 CC drug-targeting vectors for delivering anticancer drugs to cancer cells
 CC that express an apo B protein receptor, and (ii) additives for cell
 CC culture media especially as growth supplements. Non-naturally occurring,
 CC receptor-competent LDL particles do not require the complete apo B
 CC sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX

SQ Sequence 11 AA;

Query Match 84.0%; Score 42; DB 2; Length 11;
 Best Local Similarity 90.0%; Pred. No. 0.14;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDGLK 10
 |||||
 DB 2 TRLTRKRGK 11

```

RESULT 4
AAW57207
ID AAW57207 standard; peptide; 13 AA.
XX
AC AAW57207;
XX
XX 03-AUG-1998 (first entry)
XX
XX Apo B 100 binding site peptide analogue peptide B.
XX
XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
KW growth supplement; non-natural lipid particle; low density lipoprotein;
KW LDL; receptor component; apo B100 receptor site.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FH Modified-site 1
FT /note= "attached to retinoic acid"
XX
XX WO9813385-A2.
XX
XX 02-APR-1998.
XX
XX 25-SEP-1997; 97WO-GB002610.
XX
XX 27-SEP-1996; 96GB-00020153.
XX
XX (UYST ) UNIV STRATHCLYDE.
XX
XX Halbert GW, Owens MD, Baillie G;
XX
XX WPI; 1998-230637/20.
XX
XX Non-natural lipid particle comprising peptide binding to apo B protein
PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
PT that express this receptor.
XX
XX Claim 13; Fig 7; 73pp; English.
XX
XX The present sequence represents a specifically claimed Apo B 100 binding
CC site peptide analogue which can be used as a component of a non-
CC naturally occurring, receptor-competent low density lipoprotein (LDL)
CC particle of the present invention. The LDL particle comprises at least 1
CC peptide component that has at least 1 binding site for an apo B protein
CC receptor and at least 1 lipophilic substituent. Also described in the
CC invention are peptides containing an apo B binding sequence with at least
CC 70% identity with sequences: KAEYKQKHRRH (1) or TRLTRKRGGLK (2), or their
CC dimers. Non-naturally occurring, receptor-competent LDL particles are
CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
CC cancer cells that express an apo B protein receptor, and (ii) additives
CC for cell culture media especially as growth supplements. Non-naturally
CC occurring, receptor-competent LDL particles do not require the complete
CC apo B sequence, which is large and tends to aggregate, to provide binding
CC affinity to an apo B protein receptor
XX
XX Sequence 13 AA;
SQ
Query Match 84.0%; Score 42; DB 2; Length 13;
Best Local Similarity 90.0%; Pred. No. 0.17;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGGLK 10
Dy 3 TRLTRKRGGLK 12

RESULT 5
AAW41261
ID AAW41261 standard; peptide; 15 AA.
XX
AC AAW41261;
XX
XX 19-MAY-1998 (first entry)
XX
XX Apolipoprotein B-100 fragment.
XX
XX Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
KW prothrombinase complex.
XX
XX Synthetic.
XX
XX Homo sapiens.
XX
XX WO9743311-A1.
XX
XX 20-NOV-1997.
XX
XX 09-MAY-1997; 97WO-GB001255.
XX
XX 09-MAY-1996; 96GB-00009702.
XX
XX (UNLO ) ROYAL FREE HOSPITAL SCHOOL MED.
XX
XX Bruckdorfer KR, Ettelaie C;
XX
XX WPI; 1998-008798/01.
XX
XX Peptide fragments of apo:lipo:protein B-100 with anticoagulant activity -
PT used for treating or preventing coagulation, inhibiting angiogenesis,
PT cell differentiation and apoptosis.
XX
XX Disclosure; Page 22; 60pp; English.
XX
XX This sequence is an example of the peptide of the invention. It has the
CC formula (I), or their variants with one or more internal deletions.
CC insertions or substitutions, while retaining anti-coagulant properties of
CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KKNKRRHS-X2-T-22 (I) X1 = S or
CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
CC aa. Compositions containing the peptide are used for simultaneous,
CC separate or sequential treatment of cancer, particularly to prevent
CC metastatic spread. They are also used to inhibit thromboplastin-mediated
CC processes, specifically to prevent or reduce blood coagulation (e.g.
CC during or after surgery or in cases of heart attack, stroke etc.) and to
CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
CC which is active as such or as part of a 98-aa peptide, inhibits
CC activation of the prothrombinase complex; and prevents activation of
CC factor VII on the surface of thromboplastin and of platelets by thrombin.
CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
CC smaller than apoB-100, they act more quickly
XX
XX Sequence 15 AA;
SQ
Query Match 84.0%; Score 42; DB 2; Length 15;
Best Local Similarity 90.0%; Pred. No. 0.2;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGGLK 10
Dy 1 TRLTRKRGGLK 10

RESULT 6
AAW96892
ID AAW96892 standard; peptide; 15 AA.
XX
XX AAW96892;
XX
XX 22-APR-1999 (first entry)
XX
XX ApoB-100 nuclear localisation signal sequence, residues 3353-3367.
XX
XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
KW

```

KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
 OS Homo sapiens.
 XX WO9856938-A1.
 PN 17-DEC-1998.
 XX 10-JUN-1998; 98WO-US011927.
 PF 13-JUN-1997; 97US-00874807.
 PR 14-MAY-1998; 98US-00079030.
 XX (BAYU) BAYLOR COLLEGE MEDICINE.
 PA Guevara JG, Hoogveen RC, Moore JP;
 PI WPI; 1999-070331/06.
 XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.
 XX Claim 19; Fig 13D; 293pp; English.
 XX AA969878-97 represent nuclear localisation signal sequence derived from
 CC human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein
 CC component of very-low density lipoproteins (VLDL); intermediate density
 CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The
 CC present sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis
 XX Sequence 15 AA;
 SQ

Query Match 84.0%; Score 42; DB 2; Length 15;
 Best Local Similarity 90.0%; Pred. No. 0.2;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TRLTRKDGLK 10
 Db ||||| |||
 6 TRLTRKRLGLK 15
 RESULT 7
 ABJ37575
 ID ABJ37575 standard; peptide; 20 AA.
 AC ABJ37575;
 XX 10-MAY-2003 (first entry)
 DT Heparin binding peptide sequence #28.
 DE
 XX Cytostatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;
 KW cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;
 KW rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.
 XX Unidentified.
 OS WO2003007689-A2.
 XX 30-JAN-2003.
 XX 22-JUL-2002; 2002WO-US023419.
 XX

PR 20-JUL-2001; 2001US-0306726P.
 XX (ETHZ-) ETH ZURICH.
 PA (UYZU-) UNIV ZURICH.
 XX Hubbell JA, Schoenmakers R, Maynard HD;
 XX WPI; 2003-300420/29.
 DR Use of a ligand comprising of at least one sulfated or sulfonated amino
 XX acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic
 PT retinopathy and hypoxia.
 PT Disclosure; Fig 2; 79pp; English.
 XX The invention relates to a novel ligand for binding a target biomolecule,
 CC which comprises a peptide having at least one sulphated or sulphonated
 CC amino acid and at least one amino acid chosen from neutral and positively
 CC charged amino acids. The novel ligands can be used for the treatment of
 CC e.g. tumors, rheumatoid arthritis, diabetic retinopathy and hypoxia.
 CC This sequence represents a heparin binding peptide relating to the
 CC invention
 XX Sequence 20 AA;
 SQ Query Match 84.0%; Score 42; DB 6; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.27;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TRLTRKDGLK 10
 Db ||||| |||
 7 TRLTRKRLGLK 16
 RESULT 8
 AAWS7208
 ID AAWS7208 standard; peptide; 22 AA.
 XX AAWS7208;
 AC AAWS7208;
 XX 03-AUG-1998 (first entry)
 DT Apo B 100 binding site peptide analogue peptide C.
 DE Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 XX Growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FH Modified-site 1
 FT /note= "attached to retinoic acid"
 FT Modified-site 22
 FT /note= "attached to cholesterol"
 XX WO9813385-A2.
 PN 02-APR-1998.
 PD 25-SEP-1997; 97WO-GB002610.
 XX 27-SEP-1996; 96GB-00020153.
 PR (UYST) UNIV STRATHCLYDE.
 PA Halbert GW, Owens MD, Baillie G;
 PI WPI; 1998-230637/20.
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.

XX Claim 13; Fig 7; 73pp; English.
 XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKHH (1) or TRLTRKGLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 XX Sequence 22 AA;

Query Match 84.0%; Score 42; DB 2; Length 22;
 Best Local Similarity 90.0%; Pred. No. 0.29;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGLK 10
 ||||| |||
 Db 7 TRLTRKGLK 16

RESULT 9
 AAWS7209
 ID AAWS7209 standard; peptide; 22 AA.
 XX
 AC AAWS7209;
 XX
 DT 03-AUG-1998 (first entry)
 XX
 DE Apo B 100 binding site peptide analogue peptide D.
 XX
 KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 OS Synthetic.

Key Location/Qualifiers
 FT Modified-site 1 /note= "attached to retinoic acid"
 FT
 XX WO9813385-A2.
 XX
 PD 02-APR-1998.
 XX
 XX 25-SEP-1997; 97WO-GB002610.
 XX
 XX 27-SEP-1996; 96GB-00020153.
 PR
 XX (UYST) UNIV STRATHCLYDE.
 PA

XX Halbert GW, Owens MD, Baillie G;
 PI
 XX WPI; 1998-230637/20.
 DR
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 XX Claim 13; Fig 7; 73pp; English.
 PS
 XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)

CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKHH (1) or TRLTRKGLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 XX Sequence 22 AA;

Query Match 84.0%; Score 42; DB 2; Length 22;
 Best Local Similarity 90.0%; Pred. No. 0.29;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGLK 10
 ||||| |||
 Db 7 TRLTRKGLK 16

RESULT 10
 AAEL14541
 ID AAEL14541 standard; peptide; 34 AA.
 XX
 AC AAEL14541;
 XX
 DT 17-MAY-2002 (first entry)
 XX
 DE Human apoB-100 derived peptide p62.
 XX
 KW Oxidised low density lipoprotein; oxLDL; immunoassay; periaortitis;
 KW cardiovascular disease; coronary heart disease; pre-eclampsia;
 KW non-insulin-dependent diabetes; endothelial dysfunction; human; apoB-100;
 KW peptide p62.
 XX
 OS Homo sapiens.
 XX
 PN WO200206314-A2.
 XX
 XX 24-JAN-2002.
 PD
 XX 18-JUL-2001; 2001WO-GB003212.
 PF
 XX 18-JUL-2000; 2000GB-00017641.
 PR
 XX (ARKT-) ARK THERAPEUTICS LTD.
 PA
 XX Narvanen O, Yla-Herttuala S;
 PI
 XX WPI; 2002-179777/23.
 DR
 XX New peptide useful in enzyme immunoassays for detecting oxidized low
 PT density lipoprotein which is a marker of coronary heart disease and other
 PT cardiovascular diseases, has affinity for oxidized low density
 PT lipoprotein.

XX Claim 6; Page 5; 21pp; English.
 XX
 XX The invention relates to peptides having affinity for oxidised low
 CC density lipoprotein (oxLDL), in cyclised or multimeric form. The peptide
 CC is useful in an immunoassay to determine the presence, and optionally,
 CC the amount of antibodies in a sample, having affinity for oxLDL.
 CC Preferably immobilised peptide is useful for measuring the amount of
 CC autoantibodies for oxLDL in a sample, especially a serum or plasma sample
 CC from a patient for evaluating the risk of coronary heart diseases, other
 CC cardiovascular diseases, and several other disorders such as
 CC periaortitis, pre-eclampsia, non-insulin-dependent diabetes and
 CC endothelial dysfunction. The peptide of the invention is stable, can be
 CC synthesised easily without the need to isolate proteins from a patient's

CC blood, and has a long half-life. The present sequence is human apoB-100
 CC derived peptide p62 used in the invention
 XX
 SQ Sequence 34 AA;

Query Match 84.0%; Score 42; DB 5; Length 34;
 Best Local Similarity 90.0%; Pred. No. 0.47;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 0;

QY 1 TRLTRKDGK 10
 |||||
 DB 25 TRLTRKRGK 34

RESULT 11
 AAW96876
 ID AAW96876 standard; peptide; 36 AA.

XX
 AC AAW96876;

XX 22-APR-1999 (first entry)

XX Nucleic acid binding domain from apoB-100, residues 3348-3390.

XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.

XX Homo sapiens.

XX WO9856938-A1.

XX 17-DEC-1998.

XX 10-JUN-1998; 98WO-US011927.

XX 13-JUN-1997; 97US-00874807.

XX 14-MAY-1998; 98US-00079030.

XX (BAYU) BAYLOR COLLEGE MEDICINE.

XX Guevara JG, Hoogveen RC, Moore JP;

XX WPI; 1999-070331/06.

XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.

PS Claim 16; Fig 12C; 293pp; English.

XX AAW96827-77 represent nucleic acid binding domains derived from human
 CC apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein component
 CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
 CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
 CC sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis

XX Sequence 36 AA;

Query Match 84.0%; Score 42; DB 2; Length 36;
 Best Local Similarity 90.0%; Pred. No. 0.5;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDGK 10
 |||||

DB 11 TRLTRKRGK 20

RESULT 12

AAW64587

ID AAW64587 standard; peptide; 37 AA.

XX
 AC AAW64587;

XX 23-OCT-1998 (first entry)

XX Human apolipoprotein peptide fragment #1.

XX Factor V; human; detection; protein function; blood coagulation; apo;
 KW fat metabolism; Leyden mutation; deep vein thrombosis; apolipoprotein;
 KW Alzheimer's disease; 5,10-methylenetetrahydrofolate reductase; prion;
 KW hypercysteinemia; factor VII; cardiovascular disease; pathogen; virus.

XX Homo sapiens.

XX EP857973-A2.

XX 12-AUG-1998.

XX 12-JAN-1998; 98EP-00890007.

XX 13-JAN-1997; 97AT-00000044.

XX (IMMO) IMMUNO AG.

XX Moritz B, Kiessig S, Lang H, Schenk V;

XX WPI; 1998-416142/36.

XX Detecting or quantifying mutant protein in presence of wild-type protein
 PT comprises reaction with ligand used to detect mutant blood coagulation
 PT factors or apolipoproteins for diagnosing risk of thrombosis.

XX Example 2; Page 9; 18pp; German.

XX AAW64587 and AAW64588 are fragments of human apolipoprotein which are
 CC used with Factor V protein fragments in a novel method to detect the
 CC presence of a mutated protein in a sample that may also contain the
 CC corresponding wild-type protein. The method is used to detect mutations
 CC that alter protein functions (either point mutation or small insertions
 CC or deletions), particularly in proteins involved in blood coagulation or
 CC metabolism of fat. Protein functions which are specially detectable are
 CC the Leyden mutation in factor V (associated with increased risk of deep
 CC vein thrombosis), mutations in apolipoprotein (apo) genes (certain
 CC alleles of apoE indicates increased risk of developing Alzheimer's
 CC disease), thermostable 5,10-methylenetetrahydrofolate reductase
 CC (associated with hypercysteinemia and venous thrombosis) and factor VII
 CC mutations (associated with increased risk of cardiovascular disease). The
 CC method can also be applied to proteins from pathogens, e.g. viruses or
 CC prions. The method does not require complex apparatus for polymerase
 CC chain reactions, it is simple, standardisable and reliable and is
 CC particularly suited to routine screening. It also allows mutant protein
 CC in a sample to be quantified

XX Sequence 37 AA;

Query Match 84.0%; Score 42; DB 2; Length 37;
 Best Local Similarity 90.0%; Pred. No. 0.51;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDGK 10

DB 11 TRLTRKRGK 20

RESULT 13

AAW96845

ID AAW96845 standard; peptide; 51 AA.

XX AAW96845;
AC
XX
DT 22-APR-1999 (first entry)
XX
DE Nucleic acid binding domain from apob-100.
XX
KW Human apolipoprotein B-100; apob-100; very-low density lipoprotein; VLDL;
XX apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
XX non-small cell lung carcinoma; diabetes; arteriosclerosis.
XX
OS Homo sapiens.
XX
XX WO9856938-A1.
PN
XX
PD 17-DEC-1998.
XX
PF 10-JUN-1998; 98WO-US011927.
XX
XX 13-JUN-1997; 97US-00874807.
PR
XX 14-MAY-1998; 98US-00079030.
XX
PA (BAYU) BAYLOR COLLEGE MEDICINE.
XX
PI Guevara JG, Hoogvee RC, Moore JP;
XX
DR WPI; 1999-070331/06.
XX
XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
PT used for delivering nucleic acid to cells for gene therapy and antisense
PT treatment.
XX
PS Claim 16; Page 151; 293pp; English.
XX
XX AAW96827-77 represent nucleic acid binding domains derived from human
CC apolipoprotein B-100 (apob-100). Apob-100 is a major apoprotein component
CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
CC sequence can be used in the composition of the invention. The
CC specification describes a composition that comprises LDL and
CC apolipoproteins for the binding and in vivo transport of nucleic acids.
CC The composition is used to deliver nucleic acids to eukaryotic cells, in
CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
CC molecule (or ribozyme). Specifically they are used for gene therapy of
CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
CC fibrosis and arteriosclerosis
XX
SQ Sequence 51 AA;
Query Match 84.0%; Score 42; DB 2; Length 51;
Best Local Similarity 90.0%; Pred. No. 0.72; Mismatches 0; Gaps 0;
Matches 9; Conservative 0; Indels 1; Indels 0; Gaps 0;
Qy 1 TRLTRKGLK 10
Db 6 TRLTRKGLK 15
RESULT 14
ABB37687
ID ABB37687 standard; peptide; 343 AA.
XX
AC ABB37687;
XX
DT 04-FEB-2002 (first entry)
XX
DE Peptide #5193 encoded by human foetal liver single exon probe.
XX
KW Human; foetal liver; gene expression; single exon nucleic acid probe.
XX
OS Homo sapiens.
XX

PN WO200157277-A2.
XX
PD 09-AUG-2001.
XX
PF 30-JAN-2001; 2001WO-US000669.
XX
XX 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
XX 30-JUN-2000; 2000US-00608408.
PR 03-AUG-2000; 2000US-00632366.
XX 21-SEP-2000; 2000US-0234687P.
PR 27-SEP-2000; 2000US-0236359P.
XX 04-OCT-2000; 2000GB-00024263.
XX
PA (MOLE-) MOLECULAR DYNAMICS INC.
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX WPI; 2001-483447/52.
DR
XX Human genome-derived single exon nucleic acid probes useful for analyzing
PT gene expression in human fetal liver.
XX
XX Claim 27; SEQ ID NO 30322; 639pp + Sequence Listing; English.
XX
XX The invention relates to a single exon nucleic acid probe for measuring
CC human gene expression in a sample derived from human foetal liver. The
CC single exon nucleic acid probes may be used for predicting, measuring and
CC displaying gene expression in samples derived from human fetal liver. The
CC present sequence is a peptide encoded by a single exon nucleic acid probe
CC of the invention. Note: The sequence data for this patent did not form
CC part of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 343 AA;
Query Match 84.0%; Score 42; DB 4; Length 343;
Best Local Similarity 90.0%; Pred. No. 5.5;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TRLTRKGLK 10
Db 169 TRLTRKGLK 178
RESULT 15
ABG52504
ID ABG52504 standard; peptide; 343 AA.
XX
AC ABG52504;
XX
XX 25-FEB-2003 (first entry)
DT
XX Human liver peptide, SEQ ID No 31152.
DE
XX Human; liver; cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
KW hypercholesterolaemia; coronary heart disease.
XX
OS Homo sapiens.
XX
XX WO200157273-A2.
PN
XX 09-AUG-2001.
PD
XX 30-JAN-2001; 2001WO-US000664.
PF
XX 04-FEB-2000; 2000US-0180312P.
PR 26-MAY-2000; 2000US-0207456P.
XX 30-JUN-2000; 2000US-00608408.
PR 03-AUG-2000; 2000US-00632366.
XX 21-SEP-2000; 2000US-0234687P.
PR 27-SEP-2000; 2000US-0236359P.
XX 04-OCT-2000; 2000GB-00024263.
XX

XX (MOLE-) MOLECULAR DYNAMICS INC.
 XX PA Penn SC, Hanzel DK, Chen W, Rank DR;
 XX PI WPI; 2001-488998/53.
 XX DR
 XX PT Human genome-derived single exon nucleic acid probes useful for analyzing
 XX PT gene expression in human adult liver.
 XX PS Claim 27; SEQ ID NO 31152; 658pp; English.
 XX CC The invention relates to a single exon nucleic acid probe (SENP) (I) for
 CC measuring human gene expression in a sample derived from human adult
 CC liver, comprising one of 13109 defined nucleotide sequences given in the
 CC specification (or complements/ fragments). The probe hybridises at high
 CC stringency to a nucleic acid molecule expressed in the human adult liver.
 CC (I) may be used for predicting, measuring and displaying gene expression
 CC in samples derived from human adult liver. The genes identified may be
 CC involved in genetic liver diseases such as cirrhosis,
 CC hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
 CC associated with coronary heart disease. ABG47348-ABG59930 represent human
 CC liver single exon encoded peptides of the invention. Note: The sequence
 CC information for this patent does not appear in the printed specification
 CC but was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX SQ Sequence 343 AA;

Query Match 84.0%; Score 42; DB 4; Length 343;
 Best Local Similarity 90.0%; Pred. NO. 5.5;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TRLTRKGLK 10
 Db 169 TRLTRKGLK 178
 |||||
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Search completed: January 13, 2005, 01:43:00
 Job time : 79.3798 secs

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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:30:05 ; Search time 14.4262 Seconds
(without alignments)
66.696 Million cell updates/sec

Title: US-09-823-418-10
Perfect score: 50
Sequence: 1 TRLTRKDGLK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_79.*
1: PIR1.*
2: PIR2.*
3: PIR3.*
4: PIR4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|----------|--------------------|
| 1 | 42 | 84.0 | 596 | 2 S32802 | apolipoprotein B - |
| 2 | 42 | 84.0 | 4563 | 1 LPHUB | apolipoprotein B-1 |
| 3 | 38 | 76.0 | 269 | 2 C60950 | apolipoprotein B-1 |
| 4 | 38 | 76.0 | 389 | 1 DEPGA | pyruvate dehydroge |
| 5 | 38 | 76.0 | 390 | 1 DEHUPA | pyruvate dehydroge |
| 6 | 38 | 76.0 | 390 | 1 DERTPA | pyruvate dehydroge |
| 7 | 38 | 76.0 | 390 | 2 S23506 | pyruvate dehydroge |
| 8 | 38 | 76.0 | 390 | 2 S23506 | pyruvate dehydroge |
| 9 | 38 | 76.0 | 779 | 2 JH0102 | apolipoprotein B - |
| 10 | 36 | 72.0 | 275 | 2 C60950 | apolipoprotein B-1 |
| 11 | 36 | 72.0 | 394 | 2 T46858 | molybdenum cofacto |
| 12 | 35 | 70.0 | 227 | 2 A84109 | ABC transporter (A |
| 13 | 35 | 70.0 | 325 | 2 T29604 | hypothetical prote |
| 14 | 35 | 70.0 | 480 | 2 G71050 | asparagine synthas |
| 15 | 34 | 68.0 | 131 | 2 A11540 | hypothetical prote |
| 16 | 34 | 68.0 | 190 | 2 G82634 | hypothetical prote |
| 17 | 34 | 68.0 | 857 | 2 T37459 | ribonucleotide red |
| 18 | 33 | 66.0 | 453 | 2 C83043 | probable transport |
| 19 | 33 | 66.0 | 481 | 2 C82421 | conserved hypotet |
| 20 | 33 | 66.0 | 561 | 1 S34191 | sulfite reductase |
| 21 | 33 | 66.0 | 668 | 1 S74619 | hypothetical prote |
| 22 | 33 | 66.0 | 772 | 2 I50463 | protein kinase - c |
| 23 | 32 | 64.0 | 206 | 2 B97282 | ribosomal protein |
| 24 | 32 | 64.0 | 208 | 2 C82742 | transcription regu |
| 25 | 32 | 64.0 | 265 | 2 C72380 | hypothetical prote |
| 26 | 32 | 64.0 | 272 | 2 E83363 | hypothetical prote |
| 27 | 32 | 64.0 | 274 | 2 A60950 | apolipoprotein B-1 |
| 28 | 32 | 64.0 | 280 | 2 S05572 | finger protein (cl |
| 29 | 32 | 64.0 | 289 | 2 A70751 | hypothetical prote |

| | | | | | |
|----|----|------|------|----------|--------------------|
| 30 | 32 | 64.0 | 370 | 2 A49360 | pyruvate dehydroge |
| 31 | 32 | 64.0 | 388 | 1 DEHUPA | pyruvate dehydroge |
| 32 | 32 | 64.0 | 390 | 2 AG0929 | probable major tai |
| 33 | 32 | 64.0 | 390 | 2 A10836 | probable bacteriop |
| 34 | 32 | 64.0 | 473 | 2 H72393 | hypothetical prote |
| 35 | 32 | 64.0 | 555 | 2 C45868 | glycerol-3-phospha |
| 36 | 32 | 64.0 | 584 | 2 D81265 | hypothetical prote |
| 37 | 32 | 64.0 | 687 | 2 S69723 | hypothetical prote |
| 38 | 32 | 64.0 | 689 | 2 AC1408 | transcription anti |
| 39 | 32 | 64.0 | 689 | 2 AC1784 | transcription anti |
| 40 | 32 | 64.0 | 784 | 2 JH0101 | apolipoprotein B-1 |
| 41 | 32 | 64.0 | 819 | 2 S43748 | translation elonga |
| 42 | 32 | 64.0 | 833 | 2 T32289 | hypothetical prote |
| 43 | 32 | 64.0 | 1199 | 2 T18348 | probable pol polyp |
| 44 | 31 | 62.0 | 26 | 2 S51055 | ribosomal protein |
| 45 | 31 | 62.0 | 151 | 2 T28840 | hypothetical prote |

ALIGNMENTS

RESULT 1

S32802
apolipoprotein B - crab-eating macaque (fragment)
C;Species: Macaca fascicularis (crab-eating macaque)
C;Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Jul-2004
C;Accession: S32802

R;Pape, M.E.; Castle, C.K.; Murray, R.W.; Funk, G.M.; Hunt, C.E.; Marotti, K.R.; Melchior
Biochim. Biophys. Acta 1086, 326-334, 1991
A;Title: Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional r
A;Reference number: S32802; MUID:92075708; PMID:1742325
A;Accession: S32802
A;Status: preliminary
A;Molecule type: mRNA
A;Residues: 1-596 <PAP>
A;Cross-references: UNIPROT:Q28473; EMBL:X15737; NID:G38047; PIDN:CAA33755.1; PID:G9301
C;Superfamily: apolipoprotein B

Query Match 84.0% Score 42; DB 2; Length 596;
Best Local Similarity 90.0%; Pred. No. 2.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGLK 10
||| ||| |||
Db 226 TRLTRKGLK 235

RESULT 2

LPHUB
apolipoprotein B-100 precursor - human
N;Contains: apolipoprotein B-26; apolipoprotein B-48; apolipoprotein B-74
C;Species: Homo sapiens (man)
C;Date: 28-Dec-1987 #sequence_revision 28-Dec-1987 #text_change 09-Jul-2004
C;Accession: A27850; A25679; A25263; A25266; A24320; A24684; A23817; A25774; A2
4452; I61909; I59510; I39474; I394624; I37179; PS0058
R;Ludwig, B.H.; Blackhart, B.D.; Pierotti, V.R.; Caiati, L.; Fortier, C.; Knott, T.; Sc
DNA 6, 363-372, 1987
A;Title: DNA sequence of the human apolipoprotein B gene.
A;Reference number: A27850; MUID:88003974; PMID:3652907
A;Accession: A27850
A;Molecule type: DNA
A;Residues: 1-617, 'A', 619-1929, 'F', 1931-3318, 'D', 3320-3426, 'T', 3428-3431, 'Q', 3433-3731,
A;Cross-references: UNIPROT:P04114; UNIPROT:P78482; UNIPROT:P78479; UNIPROT:Q8UMNO; UNI
R;Cladaras, C.; Hadjopoulos-Cladaras, M.; Nolte, R.T.; Atkinson, D.; Zannis, V.I.
EMBO J. 5, 3495-3507, 1986
A;Title: The complete sequence and structural analysis of human apolipoprotein B-100: r
A;Reference number: A91058; MUID:87161758; PMID:3030729
A;Accession: A25679
A;Molecule type: mRNA
A;Residues: 1-11,15-2539, 'S', 2541-3823, 'R', 3825-4563 <CLA>
A;Note: 1109-Asp was also found
R;Knott, T.J.; Wallis, S.C.; Powell, L.M.; Pease, R.J.; Lusis, A.J.; Blackhart, B.; McC
Nucleic Acids Res. 14, 7501-7503, 1986

A;Title: Complete cDNA and derived protein sequence of human apolipoprotein B-100.
A;Reference number: A93639; MUID:87016385; PMID:3763409
A;Accession: A25263
A;Molecule type: mRNA
A;Residues: 1-272, 'A', 619-617, 'A', 619-1217, 'E', 1219-2091, 'V', 2093-2366, 'T', 2366-2679, 'Q',
A;Cross-references: GB:X04506; NID:G34330; PIDN:CAA28191.1; PID:G34331
R;Law, S.W.; Grant, S.M.; Higuchi, K.; Hoshattankar, A.; Lackner, K.; Lee, N.; Brewer Jr
Proc. Natl. Acad. Sci. U.S.A. 83, 8142-8146, 1986
A;Title: Human liver apolipoprotein B-100 cDNA: complete nucleic acid and derived amino
A;Reference number: A94134; MUID:87041416; PMID:3464946
A;Accession: A25267
A;Molecule type: mRNA
A;Residues: 1-617, 'A', 619-703, 'P', 705-792, 'R', 794-1270, 'S', 1272-1866, 'G', 1868-2036, 'N', 2
A;Cross-references: GB:J02610; NID:G178803; PIDN:AAA35549.1; PID:G178804
A;Note: The codons given for residues 704, 793, 1271, 2037, 2933, 3286, 3782, 4188, and
R;Chen, S.H.; Yang, C.Y.; Chen, P.F.; Setzer, D.; Tanimura, M.; Li, W.H.; Gotto Jr., A.M
J. Biol. Chem. 261, 12918-12921, 1986
A;Title: The complete cDNA and amino acid sequence of human apolipoprotein B-100.
A;Reference number: A92556; MUID:87008488; PMID:3759943
A;Accession: A25266
A;Molecule type: mRNA
A;Residues: 1-97, 'I', 99-328, 'V', 330-644, 'I', 646-918, 'P', 920-3318, 'D', 3320-3426, 'T', 3428-
A;Cross-references: GB:J02610; NID:G178803; PIDN:AAA35549.1; PID:G178804
A;Note: A total of 2366 residues were confirmed by direct sequencing of tryptic peptides
R;Protter, A.A.; Hardman, D.A.; Sato, K.Y.; Schilling, J.W.; Yamanaka, M.; Hort, Y.J.; H
Proc. Natl. Acad. Sci. U.S.A. 83, 5678-5682, 1986
A;Title: Analysis of cDNA clones encoding the entire B-26 region of human apolipoprotein
A;Reference number: A24320; MUID:86287319; PMID:3461454
A;Accession: A24320
A;Molecule type: mRNA
A;Residues: 1-97, 'I', 99-617, 'A', 619-1044, 'LA2>
A;Cross-references: GB:M12480; NID:G178791; PIDN:AAA51751.1; PID:G178792
R;Protter, A.A.; Hardman, D.A.; Schilling, J.W.; Miller, J.; Appleby, V.; Chen, G.C.; Ki
Proc. Natl. Acad. Sci. U.S.A. 83, 1467-1471, 1986
A;Title: Isolation of a cDNA clone encoding the amino-terminal region of human apolipop
A;Reference number: A94088; MUID:86149325; PMID:3513177
A;Accession: A23817
A;Molecule type: mRNA
A;Residues: 1-291, 'PRO>
A;Cross-references: GB:M12681; NID:G178797; PIDN:AAA51753.1; PID:G178798
R;Deeb, S.S.; Motulsky, A.G.; Albers, J.J.
Proc. Natl. Acad. Sci. U.S.A. 82, 4983-4986, 1985
A;Title: A partial cDNA clone for human apolipoprotein B.
A;Reference number: A25774; MUID:85270450; PMID:3860836
A;Accession: A25774
A;Molecule type: mRNA
A;Residues: 709-791, 'SSWKAASHGCHPSAGD', 810-906 <DE>
A;Cross-references: GB:X03175; NID:G178821; PIDN:AAA51759.1; PID:G178822
R;Carlsson, P.; Darnfors, C.; Olofsson, S.O.; Bjursell, G.
Gene 49, 28-51, 1986
A;Title: Analysis of the human apolipoprotein B gene: complete structure of the B-74 reg
A;Reference number: A91565; MUID:87191999; PMID:2883086
A;Accession: A26533
A;Molecule type: mRNA
A;Residues: 1282-2721, 2742-3290, 'L', 3292-3336, 'N', 3338-3948, 'P', 3950-3963, 'Y', 3965-4180,
A;Cross-references: GB:M15421; NID:G178817; PIDN:AAA51758.1; PID:G178818
R;Hardman, D.A.; Protter, A.A.; Chen, G.C.; Schilling, J.W.; Sato, K.Y.; Lau, K.; Yamana
Biochemistry 26, 5478-5486, 1987
A;Title: Structural comparison of human apolipoproteins B-48 and B-100.
A;Reference number: A29671; MUID:88050832; PMID:3676265
A;Accession: A29671
A;Molecule type: mRNA
A;Residues: 1671-2323, 'PYW', 2327-2352, 'H', 2354-2398 <HAR>
A;Cross-references: GB:M17367; NID:G178731; PIDN:AAA51741.1; PID:G178732

R;Shoulders, C.C.; Myant, N.B.; Sidoli, A.; Rodriguez, J.C.; Cortese, C.; Baralle, F.E.
Atherosclerosis 58, 277-289, 1985
A;Title: Molecular cloning of human LDL apolipoprotein B cDNA. Evidence for more than on
A;Reference number: A90084; MUID:86130855; PMID:3841481
A;Accession: A29287
A;Molecule type: mRNA
A;Residues: 3846-4238 <SHO>
R;Pittner, R.; Wagener, R.; Stoffel, W.
Biol. Chem. Hoppe-Seyler 367, 1077-1083, 1986
A;Title: Isolation, expression and characterization of a human apolipoprotein B 100-spe
A;Reference number: A25572; MUID:87076044; PMID:3024665
A;Accession: A25572
A;Molecule type: mRNA
A;Residues: 4219-4337, 'S', 4339-4563 <PFI>
A;Cross-references: GB:M36676
R;Wei, C.P.; Chen, S.H.; Yang, C.Y.; Marcel, Y.L.; Milne, R.W.; Li, W.H.; Sparrow, J.T.
Proc. Natl. Acad. Sci. U.S.A. 82, 7265-7269, 1985
A;Reference number: A24738; MUID:86042646; PMID:2932736
A;Accession: A24738
A;Molecule type: mRNA
A;Residues: 3729-3731, 'I', 3733-3875, 'A', 3877-3948, 'P', 3950-3963, 'Y', 3965-3982, 'S', 3;
A;Cross-references: GB:M12413; NID:G178735; PIDN:AAA51742.1; PID:G178736
R;Chen, S.H.; Habib, G.; Yang, C.Y.; Gu, Z.W.; Lee, B.R.; Weng, S.; Silberman, S.R.; Ca
Science 238, 363-366, 1987
A;Title: Apolipoprotein B-48 is the product of a messenger RNA with an organ-specific i
A;Reference number: A40133; MUID:88018019; PMID:3659919
A;Accession: B40133
A;Molecule type: mRNA
A;Residues: 2165-2179 <CHI>
A;Cross-references: GB:M18036; NID:G178799; PIDN:AAA51754.1; PID:G178800
A;Note: this mRNA includes the stop codon of the organ-specific mRNA for apo48
A;Accession: A40133
A;Molecule type: protein
A;Residues: 51-75; 101-110; 129-139; 158-174; 197-207; 276-287; 298-304; 306-314; 526-532; 538-5;
36; 1486-1498; 1537-1556; 1563-1572; 1601-1610; 1647-1661; 1697-1724; 1770-1781; 1859-1897; 1968
A;Note: these fragments were derived from apo48
R;Hardman, D.A.; Protter, A.A.; Schilling, J.W.; Kane, J.P.
Biochem. Biophys. Res. Commun. 149, 1214-1219, 1987
A;Title: Carboxyl terminal analysis of human B-48 protein confirms the novel mechanism i
A;Reference number: A28002; MUID:88106542; PMID:3426612
A;Accession: A28002
A;Molecule type: mRNA
A;Residues: 2129-2179, 2181-2235 <HA2>
A;Cross-references: GB:M18471
A;Experimental source: intestine
A;Note: this mRNA from intestine includes a stop codon created by RNA editing in plac
R;Mehrabian, M.; Schumaker, V.N.; Fareed, G.C.; West, R.; Johnson, D.F.; Kirchgessner, S
Nucleic Acids Res. 13, 6937-6953, 1985
A;Title: Human apolipoprotein B: identification of cDNA clones and characterization of n
A;Reference number: A24269; MUID:86041888; PMID:3903660
A;Accession: A24269
A;Molecule type: mRNA
A;Residues: 3056-3159 <MEH>
A;Cross-references: GB:X03045; NID:G28783; PIDN:CAA26850.1; PID:G929609
R;Hoshattankar, A.V.; Higuchi, K.; Law, S.W.; Meglin, N.; Brewer Jr., H.B.
Biochem. Biophys. Res. Commun. 148, 279-285, 1987
A;Title: Identification of a novel in-frame translational stop codon in human intestine
A;Reference number: A29659; MUID:88049670; PMID:2445342
A;Accession: A29659
A;Molecule type: mRNA
A;Residues: 2169-2179 <HOS>
A;Note: the sequence shown represents the carboxyl end of apolipoprotein B-48
A;Note: two RNA species, 14.1kb and 7.5kb in length, were isolated from the human intest
ch encodes the 250K apoB-48, CAA encoding 2180-Gln is substituted by the stop codon TAA,
R;Yang, C.; Kim, T.W.; Weng, S.; Lee, B.; Yang, M.; Gotto Jr., A.M.
Proc. Natl. Acad. Sci. U.S.A. 87, 5523-5527, 1990
A;Title: Isolation and characterization of sulthydryl and disulfide peptides of human at
A;Reference number: A35783; MUID:90319144; PMID:2115173
A;Contents: disulfide bonds
A;Accession: A35783
A;Molecule type: protein
A;Residues: 28-41; 76-97, 'I', 99-100; 175-193; 206-215; 239-249; 259-266; 357-399; 455-490; 512-5

A;Note: cysteines at positions 1112, 1422, 1505, 1662, 3761, 3917, and 4217 have free su
R;Lebeouf, R.C.; Miller, C.; Shively, J.E.; Schumaker, V.N.; Ballia, M.A.; Lusis, A.J.
FEBS Lett. 170, 105-108, 1984
A;Title: Human apolipoprotein B: partial amino acid sequence.
A;Reference number: A22006; MUID:84208786; PMID:6373369
A;Accession: A22006
A;Molecule type: protein
A;Residues: 873-892, 'K', 894-896 <LE1>
A;Accession: B22006
A;Molecule type: protein
A;Residues: 3113, 'L', 3115-3130, 'R', 3132-3133, 'P', 3135-3136, 'R' <LE2>
R;Blackhart, B.D.; Ludwig, E.M.; Pierotti, V.R.; Caiati, L.; Onasch, M.A.; Wallis, S.C.;
J. Biol. Chem. 261, 15364-15367, 1986
A;Title: Structure of the human apolipoprotein B gene.
A;Reference number: A92564; MUID:87057153; PMID:2946672
A;Contents: annotation; gene structure
R;Wagener, R.; Pfitzner, R.; Stoffel, W.
Biol. Chem. Hoppe-Seyler 368, 419-425, 1987
A;Title: Studies on the organization of the human apolipoprotein B 100 gene.
A;Reference number: A90715; MUID:87271140; PMID:2886136
A;Contents: annotation; gene structure
R;Weisgraber, K.H.; Rall Jr., S.C.
J. Biol. Chem. 262, 11097-11103, 1987
A;Title: Human apolipoprotein B-100 heparin-binding sites.
A;Reference number: A92605; MUID:87280197; PMID:3301850
A;Contents: annotation; heparin binding and disulfide bond
R;Dashti, N.; Lee, D.M.; Mok, T.
Biochem. Biophys. Res. Commun. 137, 493-499, 1986
A;Title: Apolipoprotein B is a calcium binding protein.
A;Reference number: A90125; MUID:86242245; PMID:3087360
A;Contents: annotation; calcium binding
R;Carlsson, P.; Olofsson, S.O.; Bondjers, G.; Wiklund, O.; Bjursell, G.
Nucleic Acids Res. 13, 8813-8826, 1985
A;Title: Molecular cloning of human apolipoprotein B cDNA.
A;Reference number: 137178; MUID:86093680; PMID:3841204
A;Accession: 137180

Query Match 84.0%; Score 42; DB 1; Length 4563;
Best Local Similarity 90.0%; Pred. No. 15;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGGLK 10
| | | | | | | |
Db 3385 TRLTRKRGGLK 3394

RESULT 3
C60950
Apolipoprotein B-100 - golden hamster (fragment)
C;Species: Mesocricetus auratus (golden hamster)
C;Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C;Accession: C60950
R;Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A;Title: A cross-species comparison of the apolipoprotein B domain that binds to the LDL
A;Reference number: A60950; MUID:90324804; PMID:2373961
A;Accession: C60950
A;Molecule type: DNA
A;Residues: 1-269 <LAW>
A;Cross-references: UNIPROT:Q60537; UNIPROT:Q60536
C;Superfamily: apolipoprotein B
C;Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;

Query Match 76.0%; Score 38; DB 2; Length 269;
Best Local Similarity 80.0%; Pred. No. 6.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGGLK 10
: | | | | | | |
Db 216 SRLTRKRGGLK 225

RESULT 4

DEPGPA
pyruvate dehydrogenase (lipoamide) (EC 1.2.4.1) alpha chain precursor - pig (fragment)
N;Alternate names: pyruvate dehydrogenase complex, E1 component alpha chain
C;Species: Sus scrofa domestica (domestic pig)
C;Date: 31-Dec-1992 #sequence_revision 31-Dec-1992 #text_change 09-Jul-2004
C;Accession: S20813
R;Sermon, K.; DeMeirleir, L.; Elpers, I.; Lissens, W.; Liebaers, I.
submitted to the EMBL Data Library, May 1990
A;Reference number: S20813
A;Accession: S20813
A;Molecule type: mRNA
A;Residues: 1-389 <SER>
A;Cross-references: UNIPROT:P29804; EMBL:X52990; NID:g1850; PIDN:CAA37180.1; PID:g1851
A;Experimental source: muscle
C;Superfamily: pyruvate dehydrogenase (lipoamide) alpha chain; thiamin pyrophosphate-b
C;Keywords: flavoprotein; heterotetramer; mitochondrion; oxidoreductase; phosphoprotein
F;1-28/Domain: transit peptide (mitochondrion) (fragment) #status predicted <TNP>
F;29-389/Product: pyruvate dehydrogenase (lipoamide) alpha chain #status predicted <NAI
F;184-233/Domain: thiamin pyrophosphate-binding domain homology <TPB>
F;231/Binding site: phosphate (Ser) (covalent) #status experimental
F;292/Binding site: phosphate (Ser) (covalent) #status experimental
F;295/Binding site: phosphate (Ser) (covalent) #status experimental

Query Match 76.0%; Score 38; DB 1; Length 389;
Best Local Similarity 80.0%; Pred. No. 8.9;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGGLK 10
| | | | | | | |
Db 53 TVLTREDGLK 62

RESULT 5
DEHUPA
pyruvate dehydrogenase (lipoamide) (EC 1.2.4.1) alpha chain precursor - human
N;Alternate names: pyruvate dehydrogenase (PDH) complex, E1 component alpha chain
C;Species: Homo sapiens (man)
C;Date: 31-Mar-1989 #sequence_revision 31-Dec-1992 #text_change 09-Jul-2004
C;Accession: JQ0770; A33905; A28275; S11715; S11716; A28398; A29577; I56309; I54262; I5
R;Koike, K.; Urata, Y.; Mateuo, S.; Koike, M.
Gene 93, 307-311, 1990
A;Title: Characterization and nucleotide sequence of the gene encoding the human pyruva
A;Reference number: JQ0770; MUID:91033044; PMID:2227443
A;Accession: JQ0770
A;Molecule type: DNA
A;Residues: 1-390 <KOI>
A;Cross-references: UNIPROT:P08559; GB:D90084; NID:g219981; PIDN:BAA14121.1; PID:g21998
R;Ho, L.; Wexler, I.D.; Liu, T.C.; Thekkumkara, T.J.; Patel, M.S.
Proc. Natl. Acad. Sci. U.S.A. 86, 5330-5334, 1989
A;Title: Characterization of cDNAs encoding human pyruvate dehydrogenase alpha-subunit.
A;Reference number: A33905; MUID:89315791; PMID:2748588
A;Accession: A33905
A;Molecule type: mRNA
A;Residues: 1-390 <HOL>
A;Cross-references: GB:M24848; NID:g190761; PIDN:AAA36533.1; PID:g190762
R;De Meirleir, L.; Mackay, N.; Wah, A.M.L.H.; Robinson, B.H.
J. Biol. Chem. 263, 1991-1995, 1988
A;Title: Isolation of a full-length complementary DNA coding for human E-1-alpha subuni
A;Reference number: A28275; MUID:88115327; PMID:2828359
A;Accession: A28275
A;Molecule type: mRNA
A;Residues: 1-348, 'P', 350-353, 'A', 355-390 <DEM1>
A;Cross-references: EMBL:J03503; NID:g189765
A;Note: the translated sequence in GenBank entry HUMPDHE1B, release 114.0, (PIDN:AAA6000
on
R;Huh, T.L.; Chi, Y.T.; Casazza, J.P.; Veech, R.L.; Song, B.J.
submitted to the EMBL Data Library, April 1990
A;Description: Identical sequences for human brain and liver pyruvate dehydrogenase E1'
A;Reference number: S11715
A;Accession: S11715
A;Molecule type: mRNA
A;Residues: 1-390 <HUI>

A;Cross-references: EMBL:X52709; NID:G35378; PIDN:CAA36933.1; PID:G35379
A;Experimental source: brain
A;Accession: S11716
A;Molecule type: mRNA
A;Residues: 1-390 <H02>
A;Cross-references: EMBL:X52710; NID:G35380; PIDN:CAA36934.1; PID:G35381
A;Experimental source: liver
A;Koike, K.; Ohta, S.; Urata, Y.; Kagawa, Y.; Koike, M.
Proc. Natl. Acad. Sci. U.S.A. 85, 41-45, 1988
A;Title: Cloning and sequencing of cDNAs encoding alpha and beta subunits of human pyruvate dehydrogenase complex
A;Reference number: A94198; MUID:88124815; PMID:3422424
A;Accession: A28398
A;Status: significant
A;Molecule type: mRNA
A;Cross-references: GB:J03575; NID:G189737; PIDN:AAA60050.1; PID:G189738
A;Note: the sequence is extensively revised in reference JQ0770; the revised sequence is R;Dahl, H.H.M.; Hunt, S.M.; Hutchison, W.M.; Brown, G.K.
J. Biol. Chem. 262, 7398-7403, 1987
A;Title: The human pyruvate dehydrogenase complex. Isolation of cDNA clones for the E1- α subunit
A;Reference number: A29577; MUID:87222349; PMID:3034892
A;Accession: A29577
A;Status: significant
A;Molecule type: mRNA
A;Cross-references: GB:M27166; NID:G488487; PIDN:AAA60051.1; PID:G387009
A;Note: the sequence reported in this reference is incorrect due to multiple frameshift
R;Ito, M.; Huq, A.H.; Naito, E.; Saijo, T.; Takeda, E.; Kuroda, Y.
J. Inherit. Metab. Dis. 15, 848-856, 1992
A;Title: Mutation of E1 alpha gene in a female patient with pyruvate dehydrogenase deficiency
A;Reference number: I56309; MUID:93188403; PMID:1338114
A;Accession: I56309
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 374-382, 'SVDOV' <ITO>
A;Cross-references: GB:S56181; NID:G299257; PIDN:AAB25674.1; PID:G299258
R;De Meirleir, L.; Lissens, W.; Vamos, E.; Liebaers, I.
Hum. Genet. 88, 649-652, 1992
A;Title: Pyruvate dehydrogenase (PDH) deficiency caused by a 21-base pair insertion mutation
A;Reference number: I54262; MUID:92201830; PMID:1551669
A;Accession: I54262
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 300-305, 'DSYRTRE', 306-309 <DEM2>
A;Cross-references: GB:S89908; NID:G247722; PIDN:AAB21863.1; PID:G247723
R;Hansen, L.L.; Horn, N.; Dahl, H.H.; Kruse, T.A.
Hum. Mol. Genet. 3, 1021-1022, 1994
A;Title: Pyruvate dehydrogenase deficiency caused by a 33 base pair duplication in the E1- α subunit
A;Reference number: I54356; MUID:95038723; PMID:7545958
A;Accession: I54356
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 300-307, 'PPHSYRTREI', 308-311 <HAN>
A;Cross-references: GB:S75842; NID:G861533; PIDN:AAB32351.1; PID:G861534
C;Comment: The phosphorylation of several serine sites by [pyruvate dehydrogenase (lipooamide)]-phosphatase complex. The enzyme is dephosphorylated by [pyruvate dehydrogenase (lipooamide)]-phosphatase
C;Genetics:
A;Gene: GDB:PDHA1
A;Cross-references: GDB:118895; OMIM:312170
A;Map position: Xp22.1-Xp22.1
A;Introns: 19/3; 23/3; 97/3; 140/1; 170/3; 201/3; 253/3; 272/3; 300/2; 336/3
C;Complex: the E1 component (pyruvate dehydrogenase) is a heterotetramer of two alpha and two beta subunits and 6 E3 component (lipooamide dehydrogenase) dimers
C;Function:
A;Description: catalyzes the reaction of pyruvate and lipooamide (on the E2 component) to form acetyl-CoA, carbon dioxide and NADH
A;Pathway: pyruvate metabolism
A;Note: thiamin pyrophosphate is a cofactor
C;Superfamily: pyruvate dehydrogenase (lipooamide) alpha chain; thiamin pyrophosphate-binding site; flavoprotein; heterotetramer; mitochondrion; oxidoreductase; phosphoprotein; F;1-29/Domain: transit peptide (mitochondrion) #status predicted <TNP>
F;30-390/Product: pyruvate dehydrogenase (lipooamide) alpha chain #status predicted <MAT>
F;185-234/Domain: thiamin pyrophosphate-binding domain homology <TNP>
F;232,293,300/Binding site: phosphate (Ser) (covalent) #status predicted

Query Match 76.0%; Score 38; DB 1; Length 390;
Best Local Similarity 80.0%; Pred. No. 8.9;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 TRLTRKDGLK 10
DB 54 TVLTREDGLK 63
RESULT 6
DERTPA
pyruvate dehydrogenase (lipooamide) (EC 1.2.4.1) alpha chain 1 precursor - rat
N;Alternate names: pyruvate dehydrogenase complex, E1 component alpha chain
C;Species: Rattus norvegicus (Norway rat)
C;Date: 31-Dec-1992 #sequence_revision 31-Dec-1992 #text_change 09-Jul-2004
C;Accession: S21553; I56503
R;Cullingford, T.E.; Phillips, I.I.; Clark, J.J.
submitted to the EMBL Data Library, June 1992
A;Reference number: S21553
A;Accession: S21553
A;Molecule type: mRNA
A;Residues: 1-390 <CULS>
A;Cross-references: UNIPROT:P26284; EMBL:Z12158; NID:G57656; PIDN:CAA78146.1; PID:G57657
A;Experimental source: liver
R;Cullingford, T.E.; Clark, J.B.; Phillips, I.R.
J. Neurochem. 62, 1682-1690, 1994
A;Title: The pyruvate dehydrogenase complex: cloning of the rat somatic E1 alpha subunit brain.
A;Reference number: I56503; MUID:94209873; PMID:8158120
A;Accession: I56503
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 1-390 <CULP>
A;Cross-references: EMBL:Z12158; NID:G57656; PIDN:CAA78146.1; PID:G57657
A;Note: in Genbank entry RRPDHYE1A, release 113.0, the source is designated as Rattus norvegicus (Norway rat)
C;Superfamily: pyruvate dehydrogenase (lipooamide) alpha chain; thiamin pyrophosphate-binding site; flavoprotein; heterotetramer; mitochondrion; oxidoreductase; phosphoprotein; F;1-29/Domain: transit peptide (mitochondrion) #status predicted <TNP>
F;30-390/Product: pyruvate dehydrogenase (lipooamide) alpha chain #status predicted <MAT>
F;185-234/Domain: thiamin pyrophosphate-binding domain homology <TPB>
F;232/Binding site: phosphate (Ser) (covalent) #status predicted
F;293/Binding site: phosphate (Ser) (covalent) #status predicted
F;300/Binding site: phosphate (Ser) (covalent) #status predicted
Query Match 76.0%; Score 38; DB 1; Length 390;
Best Local Similarity 80.0%; Pred. No. 8.9;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 1 TRLTRKDGLK 10
DB 54 TVLTREDGLK 63
RESULT 7
DERTPA
pyruvate dehydrogenase (lipooamide) (EC 1.2.4.1) alpha chain precursor - rat
N;Alternate names: pyruvate dehydrogenase complex, E1 component alpha chain
C;Species: Rattus norvegicus (Norway rat)
C;Date: 31-Dec-1992 #sequence_revision 31-Dec-1992 #text_change 09-Jul-2004
C;Accession: S15891
R;Matuda, S.; Nakano, K.; Ohta, S.; Saheki, T.; Kawanishi, Y.; Miyata, T.
Biochim. Biophys. Acta 1089, 1-7, 1991
A;Title: The alpha-ketoadid dehydrogenase complexes. Sequence similarity of rat pyruvate dehydrogenase (lipooamide) and thiamin pyrophosphate-binding domain homology
A;Reference number: S15891; MUID:91223087; PMID:2025639
A;Accession: S15891
A;Molecule type: mRNA
A;Residues: 1-390 <MAT1>
A;Cross-references: UNIPROT:P26284
C;Superfamily: pyruvate dehydrogenase (lipooamide) alpha chain; thiamin pyrophosphate-binding site; flavoprotein; heterotetramer; mitochondrion; oxidoreductase; phosphoprotein; F;1-29/Domain: transit peptide (mitochondrion) #status predicted <TNP>
F;30-390/Product: pyruvate dehydrogenase (lipooamide) alpha chain #status predicted <MAT>
F;185-234/Domain: thiamin pyrophosphate-binding domain homology <TPB>

F:232/Binding site: phosphate (Ser) (covalent) #status predicted
 F:293/Binding site: phosphate (Ser) (covalent) #status predicted
 F:300/Binding site: phosphate (Ser) (covalent) #status predicted

Query Match 76.0%; Score 38; DB 1; Length 390;
 Best Local Similarity 80.0%; Pred. No. 8.9;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10
 Db 54 TVLTREDGLK 63

RESULT 8

S23506
 pyruvate dehydrogenase (lipoamide) (EC 1.2.4.1) - mouse
 C:Species: Mus musculus (house mouse)
 C:Date: 22-Nov-1993 #sequence_revision 03-Aug-1995 #text_change 09-Jul-2004
 C:Accession: S23506
 F:Fitgerald, J.; Hutchison, W.M.; Dahl, H.H.M.
 Biochim. Biophys. Acta 1131, 83-90, 1992
 A:Title: Isolation and characterisation of the mouse pyruvate dehydrogenase E1alpha gene
 A:Reference number: S23506; MUID:92256495; PMID:1581363
 A:Accession: S23506
 A:Status: preliminary
 A:Molecule type: mRNA
 A:Residues: 1-390 <FIT>
 A:Cross-references: UNIPROT:P35486; EMBL:M76727; NID:g200276; PIDN:AAA53046.1; PID:g2002

A:Gene: pdha-1
 C:Superfamily: pyruvate dehydrogenase (lipoamide) alpha chain; thiamin pyrophosphate-bin
 C:Keywords: mitochondrion; oxidoreductase; phosphoprotein
 F:185-234/Domain: thiamin pyrophosphate-binding domain homology <TPB>

Query Match 76.0%; Score 38; DB 2; Length 390;
 Best Local Similarity 80.0%; Pred. No. 8.9;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10
 Db 54 TVLTREDGLK 63

RESULT 9

JH0102
 apolipoprotein B - golden hamster (fragment)
 C:Species: Mesocricetus auratus (golden hamster)
 C:Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
 C:Accession: JH0102
 F:Smith, T.J.
 submitted to GenBank, June 1990
 A:Reference number: A38864
 A:Accession: JH0102
 A:Molecule type: DNA
 A:Residues: 1-779 <SMI>
 A:Cross-references: UNIPROT:Q60536; GB:M35187
 A:Note: this is a revision to the sequence from reference JH0101
 R:Smith, T.J.; Hautamaa, D.; Maeda, N.
 Gene 87, 309-310, 1990

A:Title: Sequence of the putative low-density lipoprotein receptor-binding regions of ap
 A:Reference number: JH0101; MUID:90236327; PMID:2332175
 A:Contents: annotation
 A:Note: this sequence has been revised in reference A38864
 C:Genetics:

A:Gene: apob
 C:Superfamily: apolipoprotein B
 C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;
 F:435-445/Region: receptor binding
 F:646-656/Region: receptor binding

Query Match 76.0%; Score 38; DB 2; Length 779;
 Best Local Similarity 80.0%; Pred. No. 17;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10
 Db 642 SRLTRKRGK 651

RESULT 10

E60950
 apolipoprotein B-100 - chicken (fragment)
 C:Species: Gallus gallus (chicken)
 C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
 C:Accession: E60950
 R:Law, A.; Scott, J.
 J. Lipid Res. 31, 1109-1120, 1990
 A:Title: A cross-species comparison of the apolipoprotein B domain that binds to the Lp

A:Reference number: A60950; MUID:90324804; PMID:2373961
 A:Accession: E60950
 A:Molecule type: mRNA
 A:Residues: 1-275 <LAW>
 A:Cross-references: UNIPROT:Q7LZ77
 C:Superfamily: apolipoprotein B
 C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 72.0%; Score 36; DB 2; Length 275;
 Best Local Similarity 80.0%; Pred. No. 16;
 Matches 8; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10
 Db 221 TSLTRKRGK 230

RESULT 11

T46858
 molybdenum cofactor biosynthesis protein A [imported] - Rhodobacter sphaeroides
 C:Species: Rhodobacter sphaeroides
 C:Date: 17-Mar-2000 #sequence_revision 17-Mar-2000 #text_change 12-Jul-2004
 C:Accession: T46858
 R:Mackenzie, C.; Simmons, A.E.; Kaplan, S.
 Genetics 153, 525-538, 1999

A:Title: Multiple chromosomes in bacteria. The Yin and Yang of trp gene localization in
 A:Reference number: Z24108; MUID:99442363; PMID:10511537
 A:Accession: T46858
 A:Status: preliminary; translated from GB/EMBL/DBD

A:Molecule type: DNA
 A:Residues: 1-394 <MAC>
 A:Cross-references: UNIPROT:Q9ZFA5; EMBL:AF108766; NID:g4185542; PIDN:AAD09121.1; PID:g
 A:Experimental source: strain 2.4.1
 C:Genetics:
 A:Gene: moeA
 A:Map position: I
 C:Superfamily: molybdenum cofactor biosynthesis protein, MoeA type

Query Match 72.0%; Score 36; DB 2; Length 394;
 Best Local Similarity 87.5%; Pred. No. 23;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKQGL 9
 Db 341 RLTRQDGL 348

RESULT 12

A84109
 ABC transporter (ATP-binding protein) BH3673 [imported] - Bacillus halodurans (strain C
 C:Species: Bacillus halodurans
 C:Date: 01-Dec-2000 #sequence_revision 01-Dec-2000 #text_change 09-Jul-2004
 C:Accession: A84109
 R:Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hir
 Nucleic Acids Res. 28, 4317-4331, 2000
 A:Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and
 A:Reference number: A83650; MUID:20512582; PMID:11058132
 A:Accession: A84109

A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-227 <STO>
A;Cross-references: UNIPROT:Q9K6Q5; GB:AP001519; GB:BA000004; NID:g10176109; PIDN:BA073
A;Experimental source: strain C-125
C;Genetics:
A;Gene: BH3673
C;Superfamily: short-chain ATP-binding cassette proteins; ATP-binding cassette homology

Query Match 70.0%; Score 35; DB 2; Length 227;
Best Local Similarity 100.0%; Pred. No. 21;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LTRKDLG 9
|||
Db 70 LTRKDLG 76

RESULT 13

T29604
hypothetical protein ZK816.5 - Caenorhabditis elegans
C;Species: Caenorhabditis elegans
C;Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C;Accession: T29604
R;Nhan, M.; Le, T.
A;Description: The sequence of C. elegans cosmid ZK816.
A;Reference number: 220649
A;Accession: T29604
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: DNA
A;Residues: 1-325 <NHA>
A;Cross-references: UNIPROT:Q23612; EMBL:U41018; PIDN:AAA82327.1; CESP:ZK816.5
C;Genetics:
A;Gene: CESP:ZK816.5
A;Introns: 24/1; 111/3; 170/2; 228/3; 280/1

Query Match 70.0%; Score 35; DB 2; Length 325;
Best Local Similarity 60.0%; Pred. No. 30;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGLK 10
|||
Db 148 TRIMRKNMGK 157

RESULT 14

G71050
asparagine synthase (glutamine-hydrolysing) (EC 6.3.5.4) [similarity] - Pyrococcus horik
C;Species: Pyrococcus horikoshii
C;Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 09-Jul-2004
C;Accession: G71050
R;Kawarabayashi, Y.; Sawada, M.; Horikawa, H.; H aikawa, Y.; Hino, Y.; Yamamoto, S.; Sekin
M.; Ohfuku, Y.; Funahashi, T.; Tanaka, T.; Kudo, Y.; Yamazaki, J.; Kushida, N.; Oguchi
DNA Res. 5, 55-76, 1998
A;Title: Complete sequence and gene organization of the genome of a hyper-thermophilic a
A;Reference number: A71000; MUID:98344137; PMID:9679194
A;Accession: G71050
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-480 <KAW>
A;Cross-references: UNIPROT:O59829; GB:AF000005; NID:g3236132; PIDN:BAA30201.1; PID:d103
A;Experimental source: strain OT3
A;Note: this accession replaces an interim accession for a sequence replaced by GenBank
C;Genetics:
A;Gene: PH1102
A;Keywords: asparagine biosynthesis; ligase
F;2/Active site: Cys #status predicted

Query Match 70.0%; Score 35; DB 2; Length 480;
Best Local Similarity 60.0%; Pred. No. 44;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGLK 10
|||
Db 329 TRLAREDDGK 338

RESULT 15

AI1540
hypothetical protein lin0865 [imported] - Listeria innocua (strain Clip11262)
C;Species: Listeria innocua
C;Date: 27-Nov-2001 #sequence_revision 27-Nov-2001 #text_change 09-Jul-2004
C;Accession: AI1540
R;Glaser, P.; Frangoul, L.; Buchrieser, C.; Amend, A.; Baquero, F.; Berche, P.; Bloecke
.; Dominguez-Bernal, G.; Duchaud, E.; Durand, L.; Dussurget, O.; Entian, K.D.; Fsihi, H
D.; Jones, L.M.; Karst, U.
Science 294, 849-852, 2001
A;Authors: Kreft, J.; Kuhn, M.; Kunst, F.; Kurapkat, G.; Madueno, E.; Maitournan, A.; M
ok, C.; Schluter, T.; Simoes, N.; Tierrez, A.; Vazquez-Boland, J.A.; Voss, H.; Wehland
A;Title: Comparative genomics of Listeria species.
A;Reference number: AB1077; MUID:21537279; PMID:11679669
A;Accession: AI1540
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-131 <GLA>
A;Cross-references: UNIPROT:Q92DB8; GB:AL592022; PIDN:CAC96097.1; PID:g16413316; GSPDB:
A;Experimental source: strain Clip11262
C;Genetics:
A;Gene: lin0865

Query Match 68.0%; Score 34; DB 2; Length 131;
Best Local Similarity 66.7%; Pred. No. 20;
Matches 6; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGL 9
|||
Db 74 TRIQRKQGV 82

Search completed: January 13, 2005, 01:52:36
Job time : 16.4262 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 78.0328 Seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-10
Perfect score: 50
Sequence: 1 TRLTRKDGLK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : UniProt_02.*
1: uniprot_sprot.*
2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB | ID | Description |
|------------|-------|-------------|--------|----|------------|---------------------|
| 1 | 42 | 84.0 | 414 | 2 | Q7YQR5 | Q7Yqr5 aotus vocif |
| 2 | 42 | 84.0 | 596 | 2 | Q28473 | Q28473 macaca fasc |
| 3 | 42 | 84.0 | 675 | 2 | Q90VV5 | Q90vv5 fugu rubrip |
| 4 | 42 | 84.0 | 678 | 2 | P79997 | P79997 oryzias lat |
| 5 | 42 | 84.0 | 678 | 2 | Q7T041 | Q7t041 oryzias cur |
| 6 | 42 | 84.0 | 678 | 2 | Q9PW12 | Q9pw12 oryzias lat |
| 7 | 42 | 84.0 | 3262 | 2 | Q13788 | Q13788 homo sapien |
| 8 | 42 | 84.0 | 4563 | 1 | APB_HUMAN | P04114 homo sapien |
| 9 | 42 | 84.0 | 4563 | 2 | Q7Z600 | Q7z600 homo sapien |
| 10 | 38 | 76.0 | 89 | 2 | Q9N1X8 | Q9n1x8 bos taurus |
| 11 | 38 | 76.0 | 389 | 1 | ODPA_PIG | P29804 sus scrofa |
| 12 | 38 | 76.0 | 390 | 1 | ODPA_HUMAN | P08559 homo sapien |
| 13 | 38 | 76.0 | 390 | 1 | ODPA_MOUSE | P35486 mus musculus |
| 14 | 38 | 76.0 | 390 | 1 | ODPA_RAT | P36284 rattus norv |
| 15 | 38 | 76.0 | 390 | 2 | Q8HXW9 | Q8hxw9 macaca fasc |
| 16 | 38 | 76.0 | 390 | 2 | AAH07142 | Aah07142 mus muscu |
| 17 | 38 | 76.0 | 393 | 2 | Q6DGP9 | Q6dgp9 brachydanio |
| 18 | 38 | 76.0 | 421 | 2 | Q7TN68 | Q7tn68 glaucomyx v |
| 19 | 38 | 76.0 | 432 | 2 | Q7YR10 | Q7yr10 diceros bic |
| 20 | 38 | 76.0 | 436 | 2 | Q7YQW8 | Q7yqm8 nyctimene a |
| 21 | 38 | 76.0 | 438 | 2 | Q7YQW7 | Q7yqm7 pteropus hy |
| 22 | 38 | 76.0 | 438 | 2 | Q7YR04 | Q7yr04 roussetus a |
| 23 | 38 | 76.0 | 445 | 2 | Q7YR08 | Q7yr08 chaetophrac |
| 24 | 38 | 76.0 | 445 | 2 | Q7TN64 | Q7tn64 agouti paca |
| 25 | 38 | 76.0 | 445 | 2 | Q7TN71 | Q7tn71 hydrochoeru |
| 26 | 38 | 76.0 | 445 | 2 | Q7TN72 | Q7tn72 erethizon d |
| 27 | 38 | 76.0 | 780 | 2 | Q60536 | Q60536 mesocricetu |
| 28 | 38 | 76.0 | 780 | 2 | Q60537 | Q60537 mesocricetu |
| 29 | 37 | 74.0 | 446 | 2 | Q7ULCL | Q7ulcl rhodopirell |
| 30 | 37 | 74.0 | 6995 | 2 | Q96RK2 | Q96rk2 homo sapien |
| 31 | 37 | 74.0 | 22152 | 2 | Q8WXI7 | Q8wx17 homo sapien |

| | | | | | | |
|----|----|------|-----|---|------------|--------------------|
| 32 | 36 | 72.0 | 201 | 2 | O95211 | O95211 homo sapien |
| 33 | 36 | 72.0 | 202 | 1 | SECU_HUMAN | O95997 homo sapien |
| 34 | 36 | 72.0 | 202 | 2 | Cag33416 | Cag33416 homo sapi |
| 35 | 36 | 72.0 | 208 | 2 | Q8P9V7 | Q8p9v7 xanthomonas |
| 36 | 36 | 72.0 | 208 | 2 | O8PLN6 | O8pln6 xanthomonas |
| 37 | 36 | 72.0 | 275 | 2 | Q7LZ77 | Q7lzf7 gallus gall |
| 38 | 36 | 72.0 | 387 | 2 | Q7YQN2 | Q7yqn2 phalanger o |
| 39 | 36 | 72.0 | 394 | 2 | Q9ZFA5 | Q9zfz5 rhodobacter |
| 40 | 36 | 72.0 | 400 | 2 | Q7YQM9 | Q7yqm9 ornithorhyn |
| 41 | 36 | 72.0 | 405 | 2 | Q7YQNO | Q7yqno tachyglossu |
| 42 | 36 | 72.0 | 445 | 2 | Q7TN70 | Q7tn70 dinomya bra |
| 43 | 36 | 72.0 | 452 | 2 | Q73L73 | Q73l73 treponema d |
| 44 | 36 | 72.0 | 452 | 2 | AA512506 | AA512506 treponema |
| 45 | 36 | 72.0 | 669 | 2 | Q8AAW5 | Q8aaw5 bacteroides |

ALIGNMENTS

RESULT 1

Q7YQR5 PRELIMINARY; PRT; 414 AA.
AC Q7YQR5, 1
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Actus vociferans (Spix's owl monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.
OX NCBI_TaxID=57176;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548396; AAP97352.1; -.
KW Lipoprotein.
FT NON_TER 1
FT SEQUENCE 414 AA; 45955 MW; BEFA8492157E1BDE CRC64;
SQ
Query Match 84.0%; Score 42; DB 2; Length 414;
Best Local Similarity 90.0%; Pred. No. 5.8; Mismatches 1; Indels 0; Gaps 0;
Matches 9; Conservative 0;
Qy 1 TRLTRKDGLK 10
Db 258 TRLTRKGLK 267

RESULT 2

Q28473 PRELIMINARY; PRT; 596 AA.
ID Q28473, 1
AC Q28473, 1
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Apolipoprotein B (Fragment).
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecoidea; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Liver;
RC MEDLINE=92075708; PubMed=1742325;
RA Pape M.E., Castle C.K., Murray R.W., Funk G.M., Hunt C.E.,
RA Marotti K.R., Melchior G.W.;

RT "Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional regulation.";
 RL Biochim. Biophys. Acta 1086:326-334 (1991).
 RN (2)

RP SEQUENCE FROM N.A.
 RC TISSUE=Liver;

RA Murray R.;

RL Submitted (FEB-1992) to the EMBL/GenBank/DBJ databases.

DR EMBL; X15737; CAA33755.1; -.

DR PIR; S32802; S32802.

KW Lipoprotein.

FT NON_TER 596 596 1

FT NON_TER 596 596 1

SQ SEQUENCE 596 AA; 66757 MW; B13BBA74E25C3120 CRC64;

Query Match 84.0%; Score 42; DB 2; Length 596;

Best Local Similarity 90.0%; Pred. No. 8.6;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10

Db 226 TRLTRKGLK 235

RESULT 3

Q90VW5 PRELIMINARY; PRT; 675 AA.

AC Q90VW5;

DT 01-DEC-2001 (TrEMBLrel. 19, Created)

DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)

DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)

DE Soluble guanylyl cyclase alpha subunit.

GN Name=FGCS-alpha1;

OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;

OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;

OC Tetraodontidae; Tetraodontidae; Takifugu.

OC NCBI_TaxID=31033;

RN [1]

RP SEQUENCE FROM N.A.

RA Morinaga C., Yamamoto T., Moriya Y., Suzuki N.;

RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AB062171; BAB60907.1; -.

DR EMBL; AB062169; BAB60905.1; -.

DR HSSP; P30803; IAZS.

DR GO; GO:0004383; F:guanylate cyclase activity; IEA.

DR GO; GO:0016829; F:lyase activity; IEA.

DR GO; GO:0007242; P:intracellular signaling cascade; IEA.

DR InterPro; IPR001054; G_cyclase.

DR Pfam; PF00211; Guanylate_cyc; 1.

DR SMART; SM00044; CYCC; 1.

DR PROSITE; PS00452; GUANYLATE_CYCLASES_1; 1.

DR PROSITE; PS50125; GUANYLATE_CYCLASES_2; 1.

KW Lyase.

SQ SEQUENCE 675 AA; 75498 MW; E71A283DC0369601 CRC64;

Query Match 84.0%; Score 42; DB 2; Length 596;

Best Local Similarity 90.0%; Pred. No. 8.6;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10

Db 226 TRLTRKGLK 235

RESULT 3

Q90VW5 PRELIMINARY; PRT; 675 AA.

AC Q90VW5;

DT 01-DEC-2001 (TrEMBLrel. 19, Created)

DT 01-DEC-2001 (TrEMBLrel. 19, Last sequence update)

DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)

DE Soluble guanylyl cyclase alpha subunit.

GN Name=FGCS-alpha1;

OS Fugu rubripes (Japanese pufferfish) (Takifugu rubripes).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;

OC Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;

OC Tetraodontidae; Tetraodontidae; Takifugu.

OC NCBI_TaxID=31033;

RN [1]

RP SEQUENCE FROM N.A.

RA Morinaga C., Yamamoto T., Moriya Y., Suzuki N.;

RL Submitted (MAY-2001) to the EMBL/GenBank/DBJ databases.

DR EMBL; AB062171; BAB60907.1; -.

DR EMBL; AB062169; BAB60905.1; -.

DR HSSP; P30803; IAZS.

DR GO; GO:0004383; F:guanylate cyclase activity; IEA.

DR GO; GO:0016829; F:lyase activity; IEA.

DR GO; GO:0007242; P:intracellular signaling cascade; IEA.

DR InterPro; IPR001054; G_cyclase.

DR Pfam; PF00211; Guanylate_cyc; 1.

DR SMART; SM00044; CYCC; 1.

DR PROSITE; PS00452; GUANYLATE_CYCLASES_1; 1.

DR PROSITE; PS50125; GUANYLATE_CYCLASES_2; 1.

KW Lyase.

SQ SEQUENCE 675 AA; 75498 MW; E71A283DC0369601 CRC64;

Query Match 84.0%; Score 42; DB 2; Length 596;

Best Local Similarity 88.9%; Pred. No. 9.9;

Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKDGK 10

Db 315 RLTRKDGK 323

RESULT 4

P79997 PRELIMINARY; PRT; 678 AA.

AC P79997;

DT 01-MAY-1997 (TrEMBLrel. 03, Created)

DT 01-MAY-1997 (TrEMBLrel. 03, Last sequence update)

DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)

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Oy 2 RLTRKDGLK 10
Db 316 RLTRKDGLR 324

RESULT 6
Q9PMI2 PRELIMINARY; PRT; 678 AA.
ID Q9PMI2
AC Q9PMI2;
DT 01-MAY-2000 (TrEMBLrel. 13, Created)
DT 01-MAY-2000 (TrEMBLrel. 13, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Soluble guanylyl cyclase alpha subunit.
OS Euryzias latipes (Medaka fish) (Japanese ricefish).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
OC Acanthomorpha; Acanthopterygii; Percormorpha; Atherinomorpha;
OC Belontiiformes; Adrianichthyidae; Oryziinae; Oryzias.
OX NCBI_TaxID=8090;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=99303623; PubMed=10373466;
RA Mikami T., Kusakabe T., Suzuki N.;
RT "Tandem organization of medaka fish soluble guanylyl cyclase alpha
RT and beta subunit genes. Implications for coordinated transcription of
RT two subunit genes.";
RL J. Biol. Chem. 274:18567-18573(1999).
DR EMBL; AB022280; BAA76690.1; -.
DR HSSP; P30803; 1A2S.
DR GO; GO:0004383; F:guanylate cyclase activity; IEA.
DR GO; GO:0016829; F:lyase activity; IEA.
DR GO; GO:0007242; P:intracellular signaling cascade; IEA.
DR InterPro; IPR001054; G_cyclase.
DR Pfam; PF00211; Guanylate_cyc; 1.
DR SMART; SMO0044; CYC; 1.
DR PROSITE; PS00452; GUANYLATE_CYCLASES_1; 1.
DR PROSITE; PS50125; GUANYLATE_CYCLASES_2; 1.
DR KW Lyase.
SQ SEQUENCE 678 AA; 75165 MW; 66623709CBE68A5C CRC64;

Query Match 84.0%; Score 42; DB 2; Length 678;
Best Local Similarity 88.9%; Pred. No. 9.9;
Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Oy 2 RLTRKDGLK 10
Db 316 RLTRKDGLR 324

RESULT 7
Q13788 PRELIMINARY; PRT; 3262 AA.
ID Q13788
AC Q13788;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE APOB protein (Fragment).
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87191999; PubMed=2883086;
RA Carlsson P., Darnfors C., Olofsson S.O., Bjursell G.;
RT "Analysis of the human apolipoprotein B gene; complete structure of
RT the B-74 region.";
RL Gene 49:29-51(1986)
DR EMBL; M15421; ARAA51758.1; -.
DR PIR; A27850; LPHUB.
DR GO; GO:0005576; C:extracellular; NAS.
DR GO; GO:0005319; F:lipid transporter activity; NAS.
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DR GO; GO:0006869; P:lipid transport; NAS.
FT NON TER 1
SQ SEQUENCE 3262 AA; 370140 MW; 56603BC0618DD40D CRC64;

Query Match 84.0%; Score 42; DB 2; Length 3262;
Best Local Similarity 90.0%; Pred. No. 56;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Oy 1 TLTRKDGLK 10
Db 2084 TLTRKEGLK 2093

RESULT 8
APB HUMAN
ID APB HUMAN STANDARD; PRT; 4563 AA.
AC P04114; O00502; Q13787;
DT 01-NOV-1986 (Rel. 03, Created)
DT 01-NOV-1986 (Rel. 03, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Apolipoprotein B-100 precursor (Apo B-100) [Contains: Apolipoprotein
DE B-48 (APO B-48)].
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87016385; PubMed=3763409;
RA Knott T.C., Wallis S.C., Powell L.M., Pease R.J., Luis A.J.,
RA Blackhart B., McCarthy B.J., Mahley R.W., Levy-Wilson B., Scott J.;
RT "Complete cDNA and derived protein sequence of human apolipoprotein B-
RT 100.";
RL Nucleic Acids Res. 14:7501-7503(1986).
RN [2]
RP SEQUENCE FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=88003974; PubMed=3652907;
RA Ludwig E.H., Blackhart B.D., Pierotti V.R., Caiati L., Fortier C.,
RA Knott T., Scott J., Mahley R.W., Levy-Wilson B., McCarthy B.J.;
RT "DNA sequence of the human apolipoprotein B gene.";
RL DNA 6:363-372(1987).
RN [3]
RP SEQUENCE FROM N.A., AND VARIANTS ILE-98 AND GLU-4181.
RX MEDLINE=87008488; PubMed=3759943;
RA Chen S.-H., Yang C.-Y., Chen P.-F., Setzer D., Tanimura M., Li W.-H.,
RA Gotto A.M. Jr., Chan L.;
RT "The complete cDNA and amino acid sequence of human apolipoprotein B-
RT 100.";
RL J. Biol. Chem. 261:12918-12921(1986).
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=87041416; PubMed=3464946;
RA Law S.W., Grant S.M., Higuchi K., Hospattankar A.V., Lackner K.J.,
RA Lee N., Brewer H.B. Jr.;
RT "Human liver apolipoprotein B-100 cDNA: complete nucleic acid and
RT derived amino acid sequence.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:8142-8146(1986).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=87161758; PubMed=3030729;
RA Cladaras C., Hadzopoulou-Cladaras M., Nolte R.T., Atkinson D.,
RA Zannis V.I.;
RT "The complete sequence and structural analysis of human apolipoprotein
RT B-100: relationship between apoB-100 and apoB-48 forms.";
RL EMBO J. 5:3495-3507(1986).
RN [6]
RP SEQUENCE OF 709-906 FROM N.A.
RX MEDLINE=85270450; PubMed=3860836;
RA Deeb S.S., Motulsky A.G., Albers J.J.;
RT "A partial cDNA clone for human apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 82:4983-4986(1985).
RN [7]
```

RP SEQUENCE OF 3056-3159 FROM N.A.
RX MEDLINE=86041888; PubMed=3903660;
RA Mehriani M., Schumaker V.N., Fareed G.C., West R., Johnson D.F.,
RA Kirchgessner T.G., Lin H.-C., Wang X., Ma Y., Mendiaz E., Lusis A.J.;
RT "Human apolipoprotein B: identification of cDNA clones and
RT characterization of mRNA.";
RL Nucleic Acids Res. 13:6937-6953 (1985).
RN [8]
RP SEQUENCE OF 1937-2018 AND 3811-4334 FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=86093680; PubMed=3841204;
RA Carlsson P., Olofsson S.O., Bondjers G., Darnfors C., Wiklund O.,
RA Bjursell G.;
RT "Molecular cloning of human apolipoprotein B cDNA.";
RL Nucleic Acids Res. 13:8813-8826 (1985).
RN [9]
RP SEQUENCE OF 3109-4563 FROM N.A.
RX MEDLINE=85300528; PubMed=2994225;
RA Knott T.J., Rall S.C. Jr., Innerarity T.L., Jacobson S.F., Urdea M.S.,
RA Levy-Wilson B., Powell L.M., Pease R.J., Eddy R., Nakai H., Byers M.,
RA Priestley L.M., Robertson E., Rall L.B., Betsholtz C., Shows T.B.,
RA Mahley R.W., Scott J.;
RT "Human apolipoprotein B: structure of carboxyl-terminal domains, sites
RT of gene expression, and chromosomal localization.";
RL Science 230:37-43 (1985).
RN [10]
RP SEQUENCE OF 1-291 FROM N.A.
RX MEDLINE=86149325; PubMed=3513177;
RA Protter A.A., Hardman D.A., Schilling J.W., Miller J., Appleby V.,
RA Chen G.C., Kirscher S.W., McEnroe G., Kane J.P.;
RT "Isolation of a cDNA clone encoding the amino-terminal region of human
RT apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:1467-1471 (1986).
RN [11]
RP SEQUENCE OF 1-1670 FROM N.A., AND VARIANT ILE-98.
RX MEDLINE=86287319; PubMed=3461454;
RA Protter A.A., Hardman D.A., Sato K.Y., Schilling J.W., Yamanaka M.,
RA Hort J.J., Hjertild K.A., Chen G.C., Kane J.P.;
RT "Analysis of cDNA clones encoding the entire B-26 region of human
RT apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:5678-5682 (1986).
RN [12]
RP PARTIAL SEQUENCE, AND IDENTIFICATION OF APO-B48.
RX MEDLINE=86018019; PubMed=3659919;
RA Chen S.-H., Habib G., Yang C.-H., Gu Z.-W., Lee B.R., Weng S.-H.,
RA Silberman S.R., Cai S.-J., Deslypere J.P., Rosseneu M.,
RA Gotto A.M. Jr., Li W.-H., Chan L.;
RT "Apolipoprotein B-48 is the product of a messenger RNA with an organ-
RT specific in-frame stop codon.";
RL Science 238:363-366 (1987).
RN [13]
RP DOMAINS.
RX MEDLINE=87039351; PubMed=3773997;
RA Knott T.C., Pease R.J., Powell L.M., Wallis S.C., Rall S.C. Jr.,
RA Innerarity T.L., Blackhart B., Taylor W.R., Marcel Y., Milne R.,
RA Johnson D., Fuller W., Lusis A.J., McCarthy B.J., Mahley R.W.,
RA Levy-Wilson B., Scott J.;
RT "Complete protein sequence and identification of structural domains of
RT human apolipoprotein B.";
RL Nature 323:734-738 (1986).
RN [14]
RP DOMAINS.
RA Yang C.-Y., Chen S.-H., Gianturco S.H., Bradley W.A., Sparrow J.T.,
RA Tanimura M., Li W.-H., Sparrow D.A., Deloof H., Rosseneu M.,
RA Lee F.-S., Gu Z.-W., Gotto A.M. Jr., Chan L.;
RT "Sequence, structure, receptor-binding domains and internal repeats of
RT human apolipoprotein B-100.";
RL Nature 323:738-742 (1986).
RN [15]
RP CALCULUM-BINDING DATA.
RX MEDLINE=86242245; PubMed=3087360;
RA Daehli N., Lee D.M., Mok T.;
RT "Apolipoprotein B is a calcium binding protein.";
RL Biochem. Biophys. Res. Commun. 137:493-499 (1986).

RN [16]
RP PALMITOYLATION OF CYS-1112.
RX MEDLINE=20143590; PubMed=10679026;
RA Zhao Y., McCabe J.B., Vance J., Berthiaume L.G.;
RT "Palmitoylation of apolipoprotein B is required for proper
RT intracellular sorting and transport of cholesterol esters and
RT triglycerides.";
RL Mol. Biol. Cell 11:721-734 (2000).
RN [17]
RP VARIANT SER-4338.
RX MEDLINE=91071750; PubMed=1979313;
RA Navajas M., Laurent A.-M., Moreel J.-F., Ragab A., Cambou J.-P.,
RA Cuny G., Cambien F., Roizes G.;
RT "Detection by denaturing gradient gel electrophoresis of a new
RT polymorphism in the apolipoprotein B gene.";
RL Hum. Genet. 86:91-93 (1990).
RN [18]
RP VARIANT FDB GLN-3527.
RX MEDLINE=89098975; PubMed=2563166;
RA Soria L.F., Ludwig E.H., Clarke H.R.G., Vega G.L., Grundy S.M.,
RA McCarthy B.J.;
RT "Association between a specific apolipoprotein B mutation and familial
RT defective apolipoprotein B-100.";
RL Proc. Natl. Acad. Sci. U.S.A. 86:587-591 (1989).
RN [19]
RP VARIANT LEU-2739.
RX MEDLINE=91016974; PubMed=2216805;
RA Huang L.-S., Gavish D., Breslow J.L.;
RT "Sequence polymorphism in the human apoB gene at position 8344.";
RL Nucleic Acids Res. 18:5922-5922 (1990).
RN [20]
RP VARIANT FDB CYS-3558.
RX MEDLINE=95190020; PubMed=7883971;
RA Pullinger C.R., Hennessy L.K., Chatterton J.E., Liu W., Love J.A.,
RA Mendel C.M., Frost P.H., Malloy M.J., Schumaker V.N., Kane J.P.;
RT "Familial ligand-defective apolipoprotein B. Identification of a new
RT mutation that decreases LDL receptor binding affinity.";
RL J. Clin. Invest. 95:1225-1234 (1995).
RN [21]
RP VARIANTS LEU-1437; SER-1914; LYS-2566; THR-3121; ALA-3945; MET-4128
RX MEDLINE=97044521; PubMed=8889592;
RA Poirier O., Ricard S., Behague I., Souriau C., Evans A.E.,
RA Arveiler D., Marques-Vidal P., Luc G., Roizes G., Cambien F.;
RT "Detection of new variants in the apolipoprotein B (Apo B) gene by
RT PCR-SSCP.";
RL Hum. Mutat. 8:282-285 (1996).
RN [22]
RP VARIANTS FDB GLN-3527 AND CYS-3558.
RX MEDLINE=97403938; PubMed=9259199;
RA Rabes J.P., Varret M., Saint-Jore B., Erlich D., Jondeau G.,
RA Krempf M., Giraudet P., Junien C., Boileau C.;
RT "Familial ligand-defective apolipoprotein B-100: simultaneous
RT detection of the ARG3500-->GLN and ARG3531-->CYS mutations in a French
RT population.";
RL Hum. Mutat. 10:160-163 (1997).
RN [23]
RP VARIANTS SER-1914; ARG-1923; LEU-2739; THR-3427; GLN-3432
RX MEDLINE=98141125; PubMed=9490296;
RA Leren T.P., Bakken K.S., Hoel V., Hjermann I., Berg K.;
RT "Screening for mutations of the apolipoprotein B gene causing
RT hypocholesterolemia.";
RL Hum. Genet. 102:44-49 (1998).
CC -!- FUNCTION: Apolipoprotein B is a major protein constituent of
CC chylomicrons (apo B-48), LDL (apo B-100) and VLDL (apo B-100). Apo
CC B-100 functions as a recognition signal for the cellular binding
CC and internalization of LDL particles by the apoB/E receptor.
CC -!- SUBCELLULAR LOCATION: Secreted.

Query Match

Best Local Similarity 84.0%; Score 42; DB 1; Length 4563;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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QY 1 TRLTRKDGK 10
Db 3385 TRLTRKRGK 3394

RESULT 9
Q7Z600 PRELIMINARY; PRT; 4563 AA.
AC Q7Z600;
DT 01-OCT-2003 (TRENBLrel. 25, Created)
DT 01-OCT-2003 (TRENBLrel. 25, Last sequence update)
DE 01-MAR-2004 (TRENBLrel. 26, Last annotation update)
DE Apolipoprotein B (Including Ag(X) antigen).
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
RA Ahearn M.O., Kuldane S.A., Rajkumar N., Toth E.J., Yi Q.,
RA Nickerson D.A.;
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
RL EMBL; AY324608; AAF72970.1; -.
DR GO; GO:0005319; F:lipid transporter activity; IEA.
DR GO; GO:0006869; P:lipid transport; IEA.
DR InterPro; IPR009454; DUF1081.
DR InterPro; IPR001747; Lipid transprt_N.
DR Pfam; PF06448; DUF1081; 1.
DR Pfam; PF01347; Vitellogenin_N; 1.
DR SMART; SM00638; LPD_N; 1.
KW Lipoprotein.
SQ SEQUENCE 4563 AA; 515553 MW; 030B34167CEDC63C CRC64;

Query Match 84.0%; Score 42; DB 2; Length 4563;
Best Local Similarity 90.0%; Pred. No. 81;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDGK 10
Db 3385 TRLTRKRGK 3394

RESULT 10
Q9N1X8 PRELIMINARY; PRT; 89 AA.
AC Q9N1X8;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DT 01-MAR-2004 (TRENBLrel. 26, Last annotation update)
DE Pyruvate dehydrogenase E1 component alpha subunit (Fragment).
GN Name=PDHA;
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RA Polounienko A., Blecher S.;
RL Submitted (FEB-1999) to the EMBL/GenBank/DBJ databases.
RL EMBL; AF132072; AAF36517.1; -.
DR PIR; A60225; A60225.
DR HSSP; P08559; 1N14.
DR GO; GO:0016624; F:oxidoreductase activity, acting on the alde. . .; IEA.
DR GO; GO:0008152; P:metabolism; IEA.
DR InterPro; IPR001017; Dehydrogenase_E1.
DR Pfam; PF00676; E1_dh; 1.
KW Pyruvate.
NON_TER 1
SQ SEQUENCE 1 1

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FT NON TER 89
SQ SEQUENCE 89 AA; 10293 MW; B47759C2D169292B CRC64;

Query Match 76.0%; Score 38; DB 2; Length 89;
Best Local Similarity 80.0%; Pred. No. 7.3;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDGK 10
Db 45 TVLTREDGLK 54

RESULT 11
ODPA_PIG STANDARD; PRT; 389 AA.
AC P29804;
DT 01-APR-1993 (Rel. 25, Created)
DT 01-APR-1993 (Rel. 25, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Pyruvate dehydrogenase E1 component alpha subunit, somatic form,
DE mitochondrial precursor (EC 1.2.4.1) (PDHE1-A type I) (Fragment).
GN Name=PDHA1;
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
OX NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Muscle;
RX MEDLINE=90370488; PubMed=2395657;
RA Sermon K., Demeirlair L., Elpers I., Lissens W., Liebaers I.;
RT "Characterisation of a cDNA for porcine PDH-E1 alpha and comparison
RT with the human cDNA.";
RL Nucleic Acids Res. 18:4925-4925 (1990).
CC -!- FUNCTION: The pyruvate dehydrogenase complex catalyzes the overall
CC conversion of pyruvate to acetyl-CoA and CO(2). It contains
CC multiple copies of three enzymatic components: pyruvate
CC dehydrogenase [E1], dihydrolipoamide acetyltransferase [E2] and
CC lipoamide dehydrogenase [E3].
CC -!- CATALYTIC ACTIVITY: Pyruvate + [dihydrolipoalysine-residue
CC acetyltransferase] lipoylsine = [dihydrolipoalysine-residue
CC acetyltransferase] S-acetyldihydrolipoalysine + CO(2).
CC -!- COFACTOR: Thiamine pyrophosphate.
CC -!- ENZYME REGULATION: E1 activity is regulated by phosphorylation
CC (inactivation) and dephosphorylation (activation) of the alpha
CC subunit.
CC -!- SUBUNIT: Tetramer of two alpha and two beta subunits.
CC -!- SUBCELLULAR LOCATION: Mitochondrial matrix.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
CC modified and this statement is not removed. Usage by and for commercial
CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X52990; CAA37180.1; -.
CC PIR; S20813; DEFGPA.
CC HSSP; P08559; 1N14.
CC InterPro; IPR001017; Dehydrogenase_E1.
CC Pfam; PF00676; E1_dh; 1.
CC Flavoprotein; Glycolysis; Mitochondrion; Oxidoreductase;
CC Phosphorylation; Thiamine pyrophosphate; Transit peptide.
FT NON TER 1
FT TRANSIT 1
FT CHAIN 29 389 Mitochondrion (By similarity).
FT Pyruvate dehydrogenase E1 component alpha
FT subunit, somatic form.
FT MOD_RES 231 Phosphoserine (By similarity).
FT MOD_RES 292 Phosphoserine (By similarity).
FT MOD_RES 299 Phosphoserine (By similarity).
FT SEQUENCE 389 AA; 43121 MW; E9C7DF85389A9A47 CRC64;
SQ SEQUENCE 389 AA; 43121 MW; E9C7DF85389A9A47 CRC64;

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Query Match 76.0%; Score 38; DB 1; Length 389;
 Best Local Similarity 80.0%; Pred. No. 37;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDKGLK 10
 DB 53 TVLTREDGLK 62

RESULT 12

ODPA_HUMAN
 ID ODPA_HUMAN STANDARD; PRT; 390 AA.
 AC P08559; Q9NP12;
 DT 01-AUG-1988 (Rel. 08, Created)
 DT 01-MAY-1992 (Rel. 22, Last sequence update)
 DT 01-OCT-2004 (Rel. 45, Last annotation update)
 DE Pyruvate dehydrogenase E1 component alpha subunit, somatic form,
 mitochondrial precursor (EC 1.2.4.1) (PDHE1-A type I).
 GN Name=PDHA1; Synonyms=PHE1A;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Leukocyte;
 RX MEDLINE=91033044; PubMed=2227443;
 RA Koike K., Urata Y., Matsuo S., Koike M.;
 RT "Characterization and nucleotide sequence of the gene encoding the
 RT human pyruvate dehydrogenase alpha-subunit.";
 RL Gene 93:307-311(1990).
 RN [2]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain, and Liver;
 RX MEDLINE=89315791; PubMed=2748588;
 RA Ho L., Wexler I.D., Liu T.C., Thekkumkara T.J., Patel M.S.;
 RT "Characterization of cDNAs encoding human pyruvate dehydrogenase alpha
 RT subunit.";
 RL Proc. Natl. Acad. Sci. U.S.A. 86:5330-5334 (1989).
 RN [3]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Brain, and Liver;
 RX MEDLINE=89315791; PubMed=2748588;
 RA Huh T.L., Chi Y.T., Casazza J.P., Veech R.L., Song B.J.;
 RL Submitted (APR-1990) to the EMBL/GenBank/DBJ databases.
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87222349; PubMed=3034892;
 RA Dahl H.-H.M., Hunt S.M., Hutchison W.M., Brown G.K.;
 RT "The human pyruvate dehydrogenase complex. Isolation of cDNA clones
 RT for the E1 alpha subunit, sequence analysis, and characterization of
 RT the mRNA.";
 RL J. Biol. Chem. 262:7398-7403 (1987).
 RN [5]
 RP REVISIONS.
 RX MEDLINE=89308653; PubMed=2745444;
 RA Maragos C., Hutchinson W.M., Hayasaki K., Brown G.K., Dahl H.-H.M.;
 RT "Structural organization of the gene for the E1 alpha subunit of the
 RT human pyruvate dehydrogenase complex.";
 RL J. Biol. Chem. 264:12294-12298 (1989).
 RN [6]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=88115327; PubMed=2820359;
 RA de Meirleir L., MacKay N., Wah A.M.L.H., Robinson B.H.;
 RT "Isolation of a full-length complementary DNA coding for human E1
 RT alpha subunit of the pyruvate dehydrogenase complex.";
 RL J. Biol. Chem. 263:1991-1995 (1988).
 RN [7]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=88124815; PubMed=3422424;
 RA Koike K., Ohta S., Urata Y., Kagawa Y., Koike M.;
 RT "Cloning and sequencing of cDNAs encoding alpha and beta subunits of
 RT human pyruvate dehydrogenase.";
 RL Proc. Natl. Acad. Sci. U.S.A. 85:41-45 (1988).
 RN [8]

RP SEQUENCE FROM N.A.
 RC TISSUE=Muscle;
 RX MEDLINE=22388257; PubMed=12477932; DOI=10.1073/pnas.242603899;
 RA Strausberg R.L., Feingold E.A., Grouse L.H., Dege J.G.,
 RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
 RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
 RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Heien F.,
 RA Diatchenko L., Marusina K., Farmer A.F., Rubin G.M., Hong L.,
 RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
 RA Brownstein M.J., Uesdin T.B., Toshiyuki S., Carninci P., Prange C.,
 RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullighy S.J.,
 RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
 RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
 RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
 RA Fahey M., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
 RA Whiting J., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
 RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
 RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M.,
 RA Butterfield Y.S.N., Krzywinski M.I., Skalek U., Smalios D.E.,
 RA Schnerch A., Schein J.E., Jones S.J.M., Marra M.A.;
 RT "Generation and initial analysis of more than 15,000 full-length human
 RT and mouse cDNA sequences.";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903 (2002).
 RN [9]
 RP REVIEW ON VARIANTS.
 RX MEDLINE=93244853; PubMed=1301207;
 RA Dahl H.-H.M., Brown G.K., Brown R.M., Hansen L.L., Kerr D.S.,
 RA Wexler I.D., Patel M.S., de Meirleir L., Lissens W., Chun K.,
 RA McKay N., Robinson B.H.;
 RT "Mutations and polymorphisms in the pyruvate dehydrogenase E1 alpha
 RT gene.";
 RL Hum. Mutat. 1:97-102 (1992).
 RN [10]
 RP VARIANT PDHE1 DEFICIENCY LYS-313 DEL, AND VARIANT LS HIS-378.
 RX MEDLINE=91359689; PubMed=1909401;
 RA Hansen L.L., Brown G.K., Kirby D.M., Dahl H.-H.M.;
 RT "Characterization of the mutations in three patients with pyruvate
 RT dehydrogenase E1 alpha deficiency.";
 RL J. Inherit. Metab. Dis. 14:140-151 (1991).
 RN [11]
 RP VARIANT PDHE1 DEFICIENCY CYS-302.
 RX MEDLINE=93188402; PubMed=1293379;
 RA Dahl H.-H.M., Hansen L.L., Brown R.M., Danks D.M., Rogers J.G.,
 RA Brown G.K.;
 RT "X-linked pyruvate dehydrogenase E1 alpha subunit deficiency in
 RT heterozygous females: variable manifestation of the same mutation.";
 RL J. Inherit. Metab. Dis. 15:835-847 (1992).
 RN [12]
 RP VARIANT LS ALA-258.
 RX MEDLINE=93270474; PubMed=8498846;
 RA Matthews P.M., Marchington D.R., Squier M., Land J., Brown R.M.,
 RA Brown G.K.;
 RT "Molecular genetic characterization of an X-linked form of Leigh's
 RT syndrome.";
 RL Ann. Neurol. 33:652-655 (1993).
 RN [13]
 RP VARIANTS PDHE1 DEFICIENCY MET-167; THR-199; ALA-231; GLY-263 AND
 RP LEU-292.
 RX MEDLINE=93278396; PubMed=8504306;
 RA Chun K., McKay N., Petrova-Benedict R., Robinson B.H.;
 RT "Mutations in the X-linked E1 alpha subunit of pyruvate dehydrogenase
 RT leading to deficiency of the pyruvate dehydrogenase complex.";
 RL Hum. Mol. Genet. 2:449-454 (1993).
 RN [14]
 RP VARIANT LS LEU-205.
 RX MEDLINE=94258164; PubMed=8199595;
 RA Dahl H.-H.M., Brown G.K.;
 RT "Pyruvate dehydrogenase deficiency in a male caused by a point
 RT mutation (F205L) in the E1 alpha subunit.";
 RL Hum. Mutat. 3:152-155 (1994).
 RN [15]
 RP VARIANT PDHE1 DEFICIENCY GLN-263.
 RX MEDLINE=95056975; PubMed=7967473;

RA Awata H., Endo F., Tanoue A., Kitano A., Matsuda I.;
 RT "Characterization of a point mutation in the pyruvate dehydrogenase E1
 alpha gene from two boys with primary lactic acidemia.";
 RL J. Inher. Metab. Dis. 17:189-195(1994).
 [16]
 RP VARIANTS PDHE1 DEFICIENCY CYS-72; GLY-263 AND ARG-311 DEL, AND
 RP VARIANTS LS LEU-205 AND HIS-378.
 RX MEDLINE=95193781; PubMed=7887409;
 RA Chun K., Mackay N., Petrova-Benedict R., Federico A., Fois A.,
 RA Cole D.E.C., Robertson E., Robinson B.H.;
 RT "Mutations in the X-linked E1 alpha subunit of pyruvate dehydrogenase:
 exon skipping, insertion of duplicate sequence, and missense mutations
 leading to the deficiency of the pyruvate dehydrogenase complex.";
 RT Am. J. Hum. Genet. 56:558-569(1995).
 [17]
 RL VARIANTS PDHE1 DEFICIENCY PRO-10.
 RP MEDLINE=96029268; PubMed=7573035;
 RA Takakubo F., Cartwright P., Hoogenraad N., Thorburn D.R., Collins F.,
 RA Lithgow T., Dahl H.H.;
 RT "An amino acid substitution in the pyruvate dehydrogenase E1 alpha
 gene, affecting mitochondrial import of the precursor protein.";
 RL Am. J. Hum. Genet. 57:772-780(1995).
 [18]
 RP VARIANTS PDHE1 DEFICIENCY LEU-217.
 RX MEDLINE=95267751; PubMed=7757088;
 RA Henalatha S.G., Kerr D.S., Wexler I.D., Lusk M.M., Kaung M., Du Y.,
 RA Kollu M., Schelper R.L., Patel M.S.;
 RT "Pyruvate dehydrogenase complex deficiency due to a point mutation
 (P188L) within the thiamine pyrophosphate binding loop of the E1 alpha
 subunit.";
 RT Hum. Mol. Genet. 4:315-318(1995).
 [19]
 RL VARIANTS PDHE1 DEFICIENCY CYS-72; ASP-113; ARG-162; GLY-263 AND
 RP VARIANTS HIS-302.
 RX MEDLINE=96263737; PubMed=8664900;
 RA Lissens W., de Weirleir L., Seneca S., Benelli C., Marsac C.,
 RA Poll-The B.T., Briones P., Rutenbeek W., van Diggelen O., Chaigne D.,
 RA Ranaekers V., Liebaers I.;
 RT "Mutation analysis of the pyruvate dehydrogenase E1 alpha gene in
 eight patients with a pyruvate dehydrogenase complex deficiency.";
 RL Hum. Mutat. 7:46-51(1996).
 [20]
 RP VARIANTS PDHE1 DEFICIENCY VAL-210 AND ARG-311 DEL.
 RX MEDLINE=97001225; PubMed=8844217;
 RA Tripata A., Kerr D.S., Lusk M.M., Kollu M., Tan J., Patel M.S.;
 RA "Three new mutations of the pyruvate dehydrogenase alpha subunit: a
 point mutation (M181V), 3 bp deletion (-R282), and 16 bp
 insertion/frameshift (X358VS-->TVDSQ).";
 RT Hum. Mutat. 8:180-182(1996).
 [21]
 RP VARIANTS PDHE1 DEFICIENCY CYS-302 AND HIS-302.
 RX MEDLINE=98334347; PubMed=9671272;
 RA Otero L.J., Brown R.M., Brown G.K.;
 RA "Arginine 302 mutations in the pyruvate dehydrogenase E1alpha subunit
 gene: identification of further patients and in vitro demonstration of
 pathogenicity.";
 RT Hum. Mutat. 12:114-121(1998).
 CC -!- FUNCTION: The pyruvate dehydrogenase complex catalyzes the overall
 conversion of pyruvate to acetyl-CoA and CO(2). It contains
 multiple copies of three enzymatic components: pyruvate
 dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and
 lipoamide dehydrogenase (E3).
 CC -!- CATALYTIC ACTIVITY: Pyruvate + [dihydrolipoalysine-residue
 acetyltransferase] lipoylysine = [dihydrolipoalysine-residue
 acetyltransferase] S-acetyldihydrolipoalysine + CO(2).

Query Match 76.0%; Score 38; DB 1; Length 390;
 Best Local Similarity 80.0%; Pred. No. 37;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10

Db 54 TVLTREDGK 63

RESULT 13
 ODP_MOUSE STANDARD; PRT; 390 AA.
 AC P35486;
 DT 01-JUN-1994 (Rel. 29, Created)
 DT 01-JUN-1994 (Rel. 29, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DB Pyruvate dehydrogenase E1 component alpha subunit, somatic form,
 DE mitochondrial precursor (EC 1.2.4.1) (PDHE1-A type I).
 GN Name=Pdhal; Synonym=Pdha-1;
 OS Mus musculus (Mouse).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.
 OX NCBI_TaxID=10090;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=92256495; PubMed=1581363;
 RA Fitzgerald G.F., Hutchison W.M., Dahl H.H.M.;
 RT "Isolation and characterization of the mouse pyruvate dehydrogenase E1
 alpha genes";
 RL Biochim. Biophys. Acta 1131:83-90(1992).
 CC -!- FUNCTION: The pyruvate dehydrogenase complex catalyzes the overall
 conversion of pyruvate to acetyl-CoA and CO(2). It contains
 multiple copies of three enzymatic components: pyruvate
 dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and
 lipoamide dehydrogenase (E3).
 CC -!- CATALYTIC ACTIVITY: Pyruvate + [dihydrolipoalysine-residue
 acetyltransferase] lipoylysine = [dihydrolipoalysine-residue
 acetyltransferase] S-acetyldihydrolipoalysine + CO(2).
 CC -!- COFACTOR: Thiamine pyrophosphate.
 CC -!- ENZYME REGULATION: E1 activity is regulated by phosphorylation
 (inactivation) and dephosphorylation (activation) of the alpha
 subunit.
 CC -!- SUBUNIT: Tetramer of two alpha and two beta subunits.
 CC -!- SUBCELLULAR LOCATION: Mitochondrial matrix.
 CC -!- TISSUE SPECIFICITY: In all tissues, but in very low amount in
 testis.
 CC -----
 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
 between the Swiss Institute of Bioinformatics and the EMBL Outstation -
 the European Bioinformatics Institute. There are no restrictions on its
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 or send an email to license@isb-sib.ch).
 CC -----
 DR EMBL; M76727; AAA53046.1; -.
 DR PIR; S23506; S23506.
 DR HSP; P08559; INI4.
 DR MGD; MGI:197532; Pdhal.
 GO; GO:0004738; F:pyruvate dehydrogenase activity; IMP.
 DR InterPro; IPR01017; Dehydrogenase_E1.
 Pfam; PF00676; E1_dh; 1.
 DR KMW; KMW00676; E1_dh; 1.
 KW Flavoprotein; Glycolysis; Mitochondrion; Multigene family;
 KW Oxidoreductase; Phosphorylation; Thiamine pyrophosphate;
 KW Transit peptide.
 TRANSIT 1 29 Mitochondrion (By similarity).
 CHAIN 30 390 Pyruvate dehydrogenase E1 component alpha
 subunit, somatic form.
 FT MOD_RBS 232 232 Phosphoserine (By similarity).
 FT MOD_RBS 293 293 Phosphoserine (By similarity).
 FT MOD_RBS 300 300 Phosphoserine (By similarity).
 SQ SEQUENCE 390 AA; 43231 MW; 40898944CE8E0A03 CRC64;

Query Match 76.0%; Score 38; DB 1; Length 390;
 Best Local Similarity 80.0%; Pred. No. 37;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDGK 10

Db 54 TVLTREDGK 63

```

RESULT 14
ODPA_RAT          STANDARD;      PRT;    390 AA.
ID   _ODPA_RAT
AC   P26284;
DT   01-MAY-1992 (Rel. 22, Last sequence update)
DT   01-MAY-1992 (Rel. 22, Last sequence update)
DT   05-JUL-2004 (Rel. 44, Last annotation update)
DE   Pyruvate dehydrogenase E1 component alpha subunit, somatic form,
DE   mitochondrial precursor (EC 1.2.4.1) (PDHE1-A type I).
GN   Name=PDhal;
OS   Rattus norvegicus (Rat).
OC   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC   Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX   NCBI_TaxID=10116;
RN   [1]
RP   SEQUENCE FROM N.A.
RX   MEDLINE=91223087; PubMed=2025639;
RA   Matuda S., Nakano K., Ohta S., Saheki T., Kawanishi Y., Miyata T.;
RT   "The alpha-ketoacid dehydrogenase complexes. Sequence similarity of
RT   rat pyruvate dehydrogenase with Escherichia coli and Azotobacter
RT   vinelandii alpha-ketoglutarate dehydrogenase.";
RL   Biochim. Biophys. Acta 1089:1-7(1991).
RN   [2]
RP   SEQUENCE FROM N.A.
RC   STRAIN=Sprague-Dawley; TISSUE=Liver;
RX   MEDLINE=94209873; PubMed=9158120;
RA   Cullingford T.E., Clark J.B., Phillips I.R.;
RT   "The pyruvate dehydrogenase complex: cloning of the rat somatic E1
RT   alpha subunit and its coordinate expression with the mRNAs for the E1
RT   beta, E2, and E3 catalytic subunits in developing rat brain.";
RL   J. Neurochem. 62:1682-1690(1994).
CC   -!- FUNCTION: The pyruvate dehydrogenase complex catalyzes the overall
CC   conversion of pyruvate to acetyl-CoA and CO(2). It contains
CC   multiple copies of three enzymatic components: pyruvate
CC   dehydrogenase (E1), dihydrolipoamide acetyltransferase (E2) and
CC   lipoamide dehydrogenase (E3).
CC   -!- CATALYTIC ACTIVITY: Pyruvate + [dihydrolipoyllysine-residue
CC   acetyltransferase] lipoylsine = [dihydrolipoylsine-residue
CC   acetyltransferase] S-acetyldihydrolipoylsine + CO(2).
CC   -!- COFACTOR: Thiamine pyrophosphate.
CC   -!- ENZYME REGULATION: E1 activity is regulated by phosphorylation
CC   (inactivation) and dephosphorylation (activation) of the alpha
CC   subunit.
CC   -!- SUBUNIT: Tetramer of two alpha and two beta subunits.
CC   -!- SUBCELLULAR LOCATION: Mitochondrial matrix.
CC   -!- TISSUE SPECIFICITY: In all tissues, but in very low amount in
CC   testis.
-----
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-----
EMBL; Z12158; CAA78146.1; -.
DR   PIR; S15891; DERTPA.
DR   PIR; S21553; DERTP1.
DR   HSSP; P08559; 1NI4.
DR   RGD; 3286; PDhal.
DR   InterPro; IPR001017; Dehydrogenase_E1.
DR   Pfam; PF00676; E1_dh; 1.
KW   Flavoprotein; Glycolysis; Mitochondrion; Multigene family;
KW   Oxidoreductase; Phosphorylation; Thiamine pyrophosphate;
KW   Transit peptide.
FT   TRANSIT 1 29 Mitochondrion (By similarity).
FT   CHAIN 30 390 Pyruvate dehydrogenase E1 component alpha
FT   subunit, somatic form.
FT   MOD_RES 232 232 Phosphoserine (By similarity).
FT   MOD_RES 293 293 Phosphoserine (By similarity).
FT

```

```

FT   MOD_RES 300 300 Phosphoserine (By similarity).
FT   CONFLICT 10 10 R -> H (in Ref. 2).
FT   CONFLICT 126 126 N -> T (in Ref. 2).
FT   CONFLICT 129 130 HA -> LP (in Ref. 2).
FT   CONFLICT 134 134 V -> I (in Ref. 2).
SQ   SEQUENCE 390 AA; 43212 MW; 21B78A8014460DC0 CRC64;

Query Match          76.0%; Score 38; DB 1; Length 390;
Best Local Similarity 80.0%; Pred. No. 37;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY   1 TRLTRKDGK 10
Db    54 TVLTREDGK 63
      |||:||||
      |||:||||

RESULT 15
Q8HXW9          PRELIMINARY;      PRT;    390 AA.
ID   Q8HXW9;
AC   Q8HXW9;
DT   01-MAR-2003 (TrEMBLrel. 23, Created)
DT   01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT   01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE   Pyruvate dehydrogenase E1alpha.
GN   Name=PDhal;
OS   Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC   Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC   Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC   Cercopithecinae; Macaca.
OX   NCBI_TaxID=9541;
RN   [1]
RP   SEQUENCE FROM N.A.
RC   TISSUE=Brain cerebellum cortex;
RA   Kusuda J., Osada N., Hida M., Sugano S., Hashimoto K.;
RL   Submitted (APR-2002) to the EMBL/GenBank/DBJ databases.
DR   EMBL; AB083322; BAC20601.1; -.
DR   HSSP; P08559; 1NI4.
DR   GO; GO:0014624; F.oxidoreductase activity, acting on the alde. . .; IEA.
DR   GO; GO:0008152; P:metabolism; IEA.
DR   InterPro; IPR001017; Dehydrogenase_E1.
DR   Pfam; PF00676; E1_dh; 1.
KW   Pyruvate.
SQ   SEQUENCE 390 AA; 43365 MW; 420CE38364BDB33C CRC64;

Query Match          76.0%; Score 38; DB 2; Length 390;
Best Local Similarity 80.0%; Pred. No. 37;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY   1 TRLTRKDGK 10
Db    54 TVLTREDGK 63
      |||:||||
      |||:||||

Search completed: January 13, 2005, 01:51:02
Job time : 80.0328 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 77.2131 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-11

Perfect score: 49

Sequence: 1 TELTRKGLK 10

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_23Sep04:*

- 1: Geneseqp1980s:*
- 2: Geneseqp1990s:*
- 3: Geneseqp2000s:*
- 4: Geneseqp2001s:*
- 5: Geneseqp2002s:*
- 6: Geneseqp2003as:*
- 7: Geneseqp2003bs:*
- 8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------|--------------------|
| 1 | 49 | 100.0 | 10 | 2 | AAY30692 Apo-B100 |
| 2 | 46 | 93.9 | 10 | 2 | AAY30693 Apo-B100 |
| 3 | 44 | 89.8 | 11 | 2 | AAY57205 Apo B bin |
| 4 | 44 | 89.8 | 13 | 2 | AAY57207 Apo B 100 |
| 5 | 44 | 89.8 | 15 | 2 | AAY41261 Apolipop |
| 6 | 44 | 89.8 | 15 | 2 | AAY96892 ApoB-100 |
| 7 | 44 | 89.8 | 20 | 6 | ABJ37575 Heparin b |
| 8 | 44 | 89.8 | 22 | 2 | AAY57208 Apo B 100 |
| 9 | 44 | 89.8 | 22 | 2 | AAY57209 Apo B 100 |
| 10 | 44 | 89.8 | 34 | 5 | AAE14541 |
| 11 | 44 | 89.8 | 36 | 2 | AAY96876 |
| 12 | 44 | 89.8 | 37 | 2 | AAY64587 Human apo |
| 13 | 44 | 89.8 | 51 | 2 | AAY96845 Nucleic a |
| 14 | 44 | 89.8 | 343 | 4 | ABG37687 Peptide # |
| 15 | 44 | 89.8 | 343 | 4 | ABG52504 Human liv |
| 16 | 44 | 89.8 | 377 | 2 | AAY72704 Human apo |
| 17 | 44 | 89.8 | 377 | 2 | AAR34031 Sequence |
| 18 | 44 | 89.8 | 2463 | 8 | ADJ57400 Human apo |
| 19 | 44 | 89.8 | 3923 | 2 | AAY31237 Human apo |
| 20 | 44 | 89.8 | 4536 | 2 | AAY41262 Apolipop |
| 21 | 44 | 89.8 | 4536 | 2 | AAY96826 Amino aci |
| 22 | 44 | 89.8 | 4560 | 5 | AAY98981 Human apo |
| 23 | 44 | 89.8 | 4561 | 7 | ADD48677 Human pro |
| 24 | 44 | 89.8 | 4563 | 5 | AAO15893 Human apo |
| 25 | 44 | 89.8 | 4563 | 6 | ABR40253 Human ali |

| | | | | | |
|----|----|------|------|---|--------------------|
| 26 | 44 | 89.8 | 4563 | 6 | ABU79140 |
| 27 | 44 | 89.8 | 4563 | 7 | ADF43408 |
| 28 | 44 | 89.8 | 4563 | 8 | ADH18871 Human apo |
| 29 | 44 | 89.8 | 4563 | 8 | ADH18870 Human apo |
| 30 | 44 | 89.8 | 4563 | 8 | ADO33445 Human apo |
| 31 | 44 | 89.8 | 4563 | 8 | ADO33447 Human apo |
| 32 | 44 | 89.8 | 4590 | 4 | AAU33184 Novel hum |
| 33 | 41 | 83.7 | 465 | 4 | AAB92994 Human pro |
| 34 | 41 | 83.7 | 709 | 4 | AAB95320 Human pro |
| 35 | 41 | 83.7 | 719 | 7 | ADM36086 Human acy |
| 36 | 41 | 83.7 | 720 | 3 | RAY94296 Human coe |
| 37 | 41 | 83.7 | 720 | 5 | ABG61883 Prostate |
| 38 | 41 | 83.7 | 720 | 6 | ABR63871 Human fac |
| 39 | 41 | 83.7 | 720 | 7 | ADE31735 Human 266 |
| 40 | 41 | 83.7 | 720 | 7 | ADN95365 Human BEC |
| 41 | 41 | 83.7 | 720 | 8 | ADJ45487 LXR-ligan |
| 42 | 41 | 83.7 | 720 | 8 | ADP05461 Human ace |
| 43 | 40 | 81.6 | 10 | 2 | AAY30682 Apo-B100 |
| 44 | 40 | 81.6 | 10 | 2 | AAY30687 Apo-B100 |
| 45 | 39 | 79.6 | 10 | 2 | AAY30690 Apo-B100 |

ALIGNMENTS

RESULT 1
AAY30692
ID AAY30692 standard; peptide; 10 AA.
XX
AC AAY30692;
XX
DT 17-NOV-1999 (first entry)
XX
DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS Synthetic.
OS Homo sapiens.
XX
FN W09946598-A1.
XX
PD 16-SEP-1999.
XX
PF 05-MAR-1999; 99WO-US004805.
XX
PR 10-MAR-1998; 98US-0077618P.
XX
XX (REGC) UNIV CALIFORNIA.
XX
PI Innerarity TL, Boren JOS;
XX
DR WPI; 1999-551509/46.
XX
PT Identifying compounds which affect binding of low density lipoprotein
PT with proteoglycan, used for, e.g. obtaining compounds for reducing
XX atherosclerosis.
XX
PS Claim 17; Page 57; 70pp; English.
XX
CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC receptor mutations. They were created to identify compounds which
CC modulate atherosclerosis. The peptides are derived from amino acids 3358
CC to 3367 of apoB100. The method comprises detecting compounds which affect
CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC can be used for identifying compounds which disrupt LDL-PG binding
CC without inhibiting LDL receptor binding. Such compounds can be used to
CC reduce or prevent the formation of atherosclerotic lesions and prevent
CC atherosclerosis. The transgenic non-human animals and mammals which
CC express human apo-B100 can be used as an in vivo model system for the
CC study of atherosclerosis, and in vivo assay methods for identifying
CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 10 AA;

Query Match 100.0%; Score 49; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.0095;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
 | | | | |
 DB 1 TELTRKRGK 10

RESULT 2

AAAY30693
 ID AAY30693 standard; peptide; 10 AA.

AC AAY30693;

DT 17-NOV-1999 (first entry)

DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KM low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

OS Synthetic.

OS Homo sapiens.

PN W09946598-A1.

PP 16-SEP-1999.

PP 05-MAR-1999; 99WO-US004805.

PP 10-MAR-1998; 98US-007618P.

PP (REGC) UNIV CALIFORNIA.

PI Innerarity TL, Boren JOS;

PP 1999-551509/46.

PT Identifying compounds which affect binding of low density lipoprotein
 PT with proteoglycan, used for, e.g. obtaining compounds for reducing
 PT atherosclerosis.

PS Claim 17; Page 57; 70pp; English.

CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 10 AA;

Query Match 93.9%; Score 46; DB 2; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.037;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
 | | | | |
 DB 1 TELTRKRGK 10

RESULT 3

AAW57205

ID AAW57205 standard; peptide; 11 AA.

AC AAW57205;

DT 03-AUG-1998 (first entry)

DE Apo B binding site peptide 2.

KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KM growth supplement; non-natural lipid particle; low density lipoprotein;
 KM LDL; receptor component; apo B100 receptor site.

OS Synthetic.

PN W09813385-A2.

PD 02-APR-1998.

PP 25-SEP-1997; 97WO-CB002610.

PP 27-SEP-1996; 96GB-00020153.

PP (UYST) UNIV STRATHCLYDE.

PP Halbert GW, Owens MD, Baillie G;

PP WPI; 1998-230637/20.

PT Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.

PS Claim 12; Page 52; 73pp; English.

CC The present sequence represents a specifically claimed Apo B binding site
 CC peptide which can be used as a component of a non-naturally occurring,
 CC receptor-competent low density lipoprotein (LDL) particle of the present
 CC invention. The LDL particle comprises at least 1 peptide component that
 CC has at least 1 binding site for an apo B protein receptor and at least 1
 CC lipophilic substituent. Also described in the invention are peptides
 CC containing an apo B binding sequence with at least 70% identity with
 CC sequences: KASYKKNKHH (1) or TRLTRKRGK (2), or their dimers. Non-
 CC naturally occurring, receptor-competent LDL particles are useful as: (i)
 CC drug-targeting vectors for delivering anticancer drugs to cancer cells
 CC that express an apo B protein receptor, and (ii) additives for cell
 CC culture media especially as growth supplements. Non-naturally occurring,
 CC receptor-competent LDL particles do not require the complete apo B
 CC sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor

SQ Sequence 11 AA;

Query Match 89.8%; Score 44; DB 2; Length 11;
 Best Local Similarity 90.0%; Pred. No. 0.1;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
 | | | | |
 DB 2 TRLTRKRGK 11

RESULT 4
AAW57207
ID AAW57207 standard; peptide; 13 AA.
XX
AC AAW57207;
XX
DT 03-AUG-1998 (first entry)
XX
DE Apo B 100 binding site peptide analogue peptide B.
XX
KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
KW growth supplement; non-natural lipid particle; low density lipoprotein;
KW LDL; receptor component; apo B100 receptor site.
XX
OS Synthetic.
XX
FH Key Location/Qualifiers
FT Modified-site 1 /note= "attached to retinoic acid"
XX
FN W09813385-A2.
XX
PD 02-APR-1998.
XX
PF 25-SEP-1997; 97WO-GB002610.
XX
PR 27-SEP-1996; 96GB-00020153.
XX
PA (UYST) UNIV STRATHCLYDE.
XX
FI Halbert GW, Owens MD, Baillie G;
XX
DR WPI; 1998-230637/20.
XX
PT Non-natural lipid particle comprising peptide binding to apo B protein
PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
XX that express this receptor.
PS Claim 13; Fig 7; 73pp; English.
XX
CC The present sequence represents a specifically claimed apo B 100 binding
CC site peptide analogue which can be used as a component of a non-
CC naturally occurring, receptor-competent low density lipoprotein (LDL)
CC particle of the present invention. The LDL particle comprises at least 1
CC peptide component that has at least 1 binding site for an apo B protein
CC receptor and at least 1 lipophilic substituent. Also described in the
CC invention are peptides containing an apo B binding sequence with at least
CC 70% identity with sequences: KAAYKKNKRRH (1) or TRLTRKRGGLK (2), or their
CC dimers. Non-naturally occurring, receptor-competent LDL particles are
CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
CC cancer cells that express an apo B protein receptor, and (ii) additives
CC for cell culture media especially as growth supplements. Non-naturally
CC occurring, receptor-competent LDL particles do not require the complete
CC apo B sequence, which is large and tends to aggregate, to provide binding
CC affinity to an apo B protein receptor
XX
SQ Sequence 13 AA;
Query Match 89.8%; Score 44; DB 2; Length 13;
Best Local Similarity 90.0%; Pred. No. 0.12;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TELTRKRGGLK 10
Db 3 TRLTRKRGGLK 12
RESULT 5
AAW41261
ID AAW41261 standard; peptide; 15 AA.
XX
AC AAW41261;

XX 19-MAY-1998 (first entry)
DT
DE Apolipoprotein B-100 fragment.
XX
KW Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
KW prothrombinase complex.
XX
OS Synthetic.
OS Homo sapiens.
XX W09743311-A1.
PN
XX 20-NOV-1997.
PD
XX 09-MAY-1997; 97WO-GB001255.
PF
XX 09-MAY-1996; 96GB-00009702.
PR
XX (UNLO) ROYAL FREE HOSPITAL SCHOOL MED.
PA
XX Bruckdorfer KR, Eteleaie C;
XX
XX WPI; 1998-008798/01.
DR
XX Peptide fragments of apo-lipo:protein B-100 with anticoagulant activity -
PT used for treating or preventing coagulation, inhibiting angiogenesis,
PT cell differentiation and apoptosis.
XX
XX Disclosure; Page 22; 60pp; English.
XX
CC This sequence is an example of the peptide of the invention. It has the
CC formula (I), or their variants with one or more internal deletions,
CC insertions or substitutions, while retaining anti-coagulant properties of
CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KKNKRRHS-X2-T-Z2 (I) X1 = S or
CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
CC aa. Compositions containing the peptide are used for simultaneous,
CC separate or sequential treatment of cancer, particularly to prevent
CC metastatic spread. They are also used to inhibit thromboplastin-mediated
CC processes, specifically to prevent or reduce blood coagulation (e.g.
CC during or after surgery or in cases of heart attack, stroke etc.) and to
CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
CC which is active as such or as part of a 98-aa peptide, inhibits
CC activation of the prothrombinase complex; and prevents activation of
CC factor VII on the surface of thromboplastin and of platelets by thrombin.
CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
CC smaller than apoB-100, they act more quickly
XX
SQ Sequence 15 AA;
Query Match 89.8%; Score 44; DB 2; Length 15;
Best Local Similarity 90.0%; Pred. No. 0.14;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TELTRKRGGLK 10
Db 1 TRLTRKRGGLK 10
RESULT 6
AAW96892
ID AAW96892 standard; peptide; 15 AA.
XX
XX AAW96892;
AC
XX
XX 22-APR-1999 (first entry)
DT
XX ApoB-100 nuclear localisation signal sequence, residues 3353-3367.
DE
XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
KW

KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
 OS Homo sapiens.
 XX WO9856938-A1.
 PN 17-DEC-1998.
 XX 10-JUN-1998; 98WO-US011927.
 PF 13-JUN-1997; 97US-00874807.
 PR 14-MAY-1998; 98US-00079030.
 XX (BAYU) BAYLOR COLLEGE MEDICINE.
 PA Guevara JG, Hoogveen RC, Moore JP;
 PI WPI; 1999-070331/06.
 DR Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.
 XX Claim 19; Fig 13D; 293pp; English.
 PS AAW96878-97 represent nuclear localisation signal sequence derived from
 XX human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein
 CC component of very-low density lipoproteins (VLDL), intermediate density
 CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The
 CC present sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis
 XX SQ Sequence 15 AA;
 Query Match 89.8%; Score 44; DB 2; Length 15;
 Best Local Similarity 90.0%; Pred. No. 0.14;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TELTRKRGGLK 10
 | | | | |
 Db 6 TRLTRKRGGLK 15
 RESULT 7
 ABJ37575
 ID ABJ37575 standard; peptide; 20 AA.
 XX AC ABJ37575;
 XX 10-MAY-2003 (first entry)
 DT Heparin binding peptide sequence #28.
 DE Cytostatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;
 KW cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;
 KW rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.
 XX Unidentified.
 OS WO2003007689-A2.
 XX PN 30-JAN-2003.
 PD 22-JUL-2002; 2002WO-US023419.
 XX PF Non-natural lipid particle comprising peptide binding to apo B protein
 XX receptor - useful as, e.g. vector for delivering drugs to cancer cells
 XX that express this receptor.

PR 20-JUL-2001; 2001US-0306726P.
 XX (ETHZ-) ETH ZUERICH.
 PA (UYZU-) UNIV ZURICH.
 XX Hubbell JA, Schoenmakers R, Maynard HD;
 PI WPI; 2003-300420/29.
 DR Use of a ligand comprising of at least one sulfated or sulfonated amino
 XX acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic
 PT retinopathy and hypoxia.
 PT Disclosure; Fig 2; 79pp; English.
 PS The invention relates to a novel ligand for binding a target biomolecule,
 XX which comprises a peptide having at least one sulphated or sulphonated
 CC amino acid and at least one amino acid chosen from neutral and positively
 CC charged amino acids. The novel ligands can be used for the treatment of
 CC e.g. tumors, rheumatoid arthritis, diabetic retinopathy and hypoxia.
 CC This sequence represents a heparin binding peptide relating to the
 CC invention
 XX SQ Sequence 20 AA;
 Query Match 89.8%; Score 44; DB 6; Length 20;
 Best Local Similarity 90.0%; Pred. No. 0.18;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TELTRKRGGLK 10
 | | | | |
 Db 7 TRLTRKRGGLK 16
 RESULT 8
 AAW57208
 ID AAW57208 standard; peptide; 22 AA.
 XX AC AAW57208;
 XX 03-AUG-1998 (first entry)
 DT Apo B 100 binding site peptide analogue peptide C.
 DE Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 XX growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 KW Synthetic.
 XX OS
 XX Key Location/Qualifiers
 FH Modified-site 1
 FT Modified-site /note= "attached to retinoic acid"
 FT Modified-site 22
 FT /note= "attached to cholesterol"
 XX WO9813385-A2.
 PN 02-APR-1998.
 PD 25-SEP-1997; 97WO-GB002610.
 PF 27-SEP-1996; 96GB-00020153.
 PR (UYST) UNIV STRATHCLYDE.
 PA Halbert GW, Owens MD, Baillie G;
 XX WPI; 1998-230637/20.
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 XX receptor - useful as, e.g. vector for delivering drugs to cancer cells
 XX that express this receptor.

XX Claim 13; Fig 7; 73pp; English.

XX The present sequence represents a specifically claimed Apo B 100 binding

CC site peptide analogue which can be used as a component of a non-

CC naturally occurring, receptor-competent low density lipoprotein (LDL)

CC particle of the present invention. The LDL particle comprises at least 1

CC peptide component that has at least 1 binding site for an apo B protein

CC receptor and at least 1 lipophilic substituent. Also described in the

CC invention are peptides containing an apo B binding sequence with at least

CC 70% identity with sequences: KAEYKKNKHH (1) or TRLTRKGLK (2), or their

CC dimers. Non-naturally occurring, receptor-competent LDL particles are

CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to

CC cancer cells that express an apo B protein receptor, and (ii) additives

CC for cell culture media especially as growth supplements. Non-naturally

CC occurring, receptor-competent LDL particles do not require the complete

CC apo B sequence, which is large and tends to aggregate, to provide binding

CC affinity to an apo B protein receptor

XX SQ Sequence 22 AA;

Query Match 89.8%; Score 44; DB 2; Length 22;

Best Local Similarity 90.0%; Pred. No. 0.2;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10

Db 7 TRLTRKGLK 16

RESULT 9

AAW57209

ID AAW57209 standard; peptide; 22 AA.

AC AAW57209;

XX 03-AUG-1998 (first entry)

DE Apo B 100 binding site peptide analogue peptide D.

XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;

KW growth supplement; non-natural lipid particle; low density lipoprotein;

KW LDL; receptor component; apo B100 receptor site.

XX Synthetic.

XX Key Location/Qualifiers

FT Modified-site 1 /note= "attached to retinoic acid"

XX WO9813385-A2.

XX 02-APR-1998.

XX 25-SEP-1997; 97WO-GB002610.

XX 27-SEP-1996; 96GB-00020153.

XX (UYST) UNIV STRATHCLYDE.

XX Halbert GW, Owens MD, Baillie G;

XX WPI; 1998-230637/20.

XX Non-natural lipid particle comprising peptide binding to apo B protein

FT receptor - useful as, e.g. vector for delivering drugs to cancer cells

PT that express this receptor.

XX Claim 13; Fig 7; 73pp; English.

XX The present sequence represents a specifically claimed Apo B 100 binding

CC site peptide analogue which can be used as a component of a non-

CC naturally occurring, receptor-competent low density lipoprotein (LDL)

CC particle of the present invention. The LDL particle comprises at least 1

CC peptide component that has at least 1 binding site for an apo B protein

CC receptor and at least 1 lipophilic substituent. Also described in the

CC invention are peptides containing an apo B binding sequence with at least

CC 70% identity with sequences: KAEYKKNKHH (1) or TRLTRKGLK (2), or their

CC dimers. Non-naturally occurring, receptor-competent LDL particles are

CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to

CC cancer cells that express an apo B protein receptor, and (ii) additives

CC for cell culture media especially as growth supplements. Non-naturally

CC occurring, receptor-competent LDL particles do not require the complete

CC apo B sequence, which is large and tends to aggregate, to provide binding

CC affinity to an apo B protein receptor

XX SQ Sequence 22 AA;

Query Match 89.8%; Score 44; DB 2; Length 22;

Best Local Similarity 90.0%; Pred. No. 0.2;

Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10

Db 7 TRLTRKGLK 16

RESULT 10

AAE14541

ID AAE14541 standard; peptide; 34 AA.

XX AAE14541;

XX 17-MAY-2002 (first entry)

XX Human apoB-100 derived peptide p62.

XX Oxidised low density lipoprotein; oxLDL; immunoassay; periaortitis;

KW cardiovascular disease; coronary heart disease; pre-eclampsia;

KW non-insulin-dependent diabetes; endothelial dysfunction; human; apoB-100;

XX peptide p62.

XX Homo sapiens.

XX WO200206314-A2.

XX 24-JAN-2002.

XX 18-JUL-2001; 2001WO-GB003212.

XX 18-JUL-2000; 2000GB-00017641.

XX (ARKT-) ARK THERAPEUTICS LTD.

XX Narvanen O, Yla-Herttuala S;

XX WPI; 2002-179777/23.

XX New peptide useful in enzyme immunoassays for detecting oxidized low

XX density lipoprotein which is a marker of coronary heart disease and other

XX cardiovascular diseases, has affinity for oxidized low density

XX lipoprotein.

XX Claim 6; Page 5; 21pp; English.

XX The invention relates to peptides having affinity for oxidised low

XX density lipoprotein (oxLDL), in cyclised or multimeric form. The peptide

XX is useful in an immunoassay to determine the presence, and optionally,

XX the amount of antibodies in a sample, having affinity for oxLDL.

XX Preferably immobilised peptide is useful for measuring the amount of

XX autoantibodies for oxLDL in a sample, especially a serum or plasma sample

XX from a patient for evaluating the risk of coronary heart diseases, other

XX cardiovascular diseases, and several other disorders such as

XX periaortitis, pre-eclampsia, non-insulin-dependent diabetes and

XX endothelial dysfunction. The peptide of the invention is stable, can be

XX synthesised easily without the need to isolate proteins from a patient's

CC blood, and has a long half-life. The present sequence is human apoB-100
 CC derived peptide p62 used in the invention
 XX
 SQ

Sequence 34 AA;

Query Match 89.8%; Score 44; DB 5; Length 34;
 Best Local Similarity 90.0%; Pred. No. 0.31;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLGLK 10
 | | | | | | | | | |
 Db 25 TELTRKRLGLK 34

RESULT 11
 AAW96876
 ID AAW96876 standard; peptide; 36 AA.

XX AAW96876;

AC
 XX
 DT 22-APR-1999 (first entry)

DE Nucleic acid binding domain from apoB-100, residues 3348-3390.

XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.

XX Homo sapiens.

OS
 XX
 PN WO9856938-A1.

XX 17-DEC-1998.

XX 10-JUN-1998; 98WO-US011927.

XX 13-JUN-1997; 97US-00874807.

PR 14-MAY-1998; 98US-00079030.

XX (BAYU) BAYLOR COLLEGE MEDICINE.

XX Guevara JG, Hoogveen RC, Moore JP;

XX WPI; 1999-070331/06.

PT Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.

XX Claim 16; Fig 12C; 293pp; English.

XX AAW96827-77 represent nucleic acid binding domains derived from human
 CC apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein component
 CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
 CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
 CC sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis

SQ Sequence 36 AA;

Query Match 89.8%; Score 44; DB 2; Length 36;
 Best Local Similarity 90.0%; Pred. No. 0.33;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLGLK 10
 | | | | | | | | | |

Db 11 TELTRKRLGLK 20

RESULT 12

AAW64587

ID AAW64587 standard; peptide; 37 AA.

XX AAW64587;

AC
 XX 23-OCT-1998 (first entry)

XX Human apolipoprotein peptide fragment #1.

XX Factor V; human; detection; protein function; blood coagulation; apo;
 KW fat metabolism; Leyden mutation; deep vein thrombosis; apolipoprotein;
 KW Alzheimer's disease; 5,10-methylenetetrahydrofolate reductase; prion;
 KW hypercystinaemia; factor VII; cardiovascular disease; pathogen; virus.

XX Homo sapiens.

OS
 XX
 PN EP857973-A2.

XX 12-AUG-1998.

XX 12-JAN-1998; 98EP-00890007.

XX 13-JAN-1997; 97AT-00000044.

XX (IMMO) IMMUNO AG.

XX Moritz B, Kiessig S, Lang H, Schenk V;

XX WPI; 1998-416142/36.

XX Detecting or quantifying mutant protein in presence of wild-type protein
 PT comprises reaction with ligand - used to detect mutant blood coagulation
 PT factors or apolipoproteins for diagnosing risk of thrombosis.

XX Example 2; Page 9; 18pp; German.

XX AAW64587 and AAW64588 are fragments of human apolipoprotein which are
 CC used with Factor V protein fragments in a novel method to detect the
 CC presence of a mutated protein in a sample that may also contain the
 CC corresponding wild-type protein. The method is used to detect mutations
 CC that alter protein functions (either point mutation or small insertions
 CC or deletions), particularly in proteins involved in blood coagulation or
 CC metabolism of fat. Protein functions which are specially detectable are
 CC the Leyden mutation in factor V (associated with increased risk of deep
 CC vein thrombosis), mutations in apolipoprotein (apo) genes (certain
 CC alleles of apoE indicates increased risk of developing Alzheimer's
 CC disease), thermostable 5,10-methylenetetrahydrofolate reductase
 CC (associated with hypercystinaemia and venous thrombosis) and factor VII
 CC mutations (associated with increased risk of cardiovascular disease). The
 CC method can also be applied to proteins from pathogens, e.g. viruses or
 CC prions. The method does not require complex apparatus for polymerase
 CC chain reactions, it is simple, standardisable and reliable and is
 CC particularly suited to routine screening. It also allows mutant protein
 CC in a sample to be quantified

SQ Sequence 37 AA;

Query Match 89.8%; Score 44; DB 2; Length 37;
 Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLGLK 10
 | | | | | | | | | |
 Db 11 TELTRKRLGLK 20

RESULT 13

AAW96845

ID AAW96845 standard; peptide; 51 AA.

XX AAW96845;
AC 22-APR-1999 (first entry)
XX Nucleic acid binding domain from apob-100.
XX Human apolipoprotein B-100; apob-100; very-low density lipoprotein; VLDL;
KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
XX Homo sapiens.
XX WO9856938-A1.
XX 17-DEC-1998.
XX 10-JUN-1998; 98WO-US011927.
XX 13-JUN-1997; 97US-00874807.
XX 14-MAY-1998; 98US-00079030.
XX (BAYU) BAYLOR COLLEGE MEDICINE.
XX Guevara JG, Hoogveen RC, Moore JP;
XX WPI; 1999-070331/06.
XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
PT used for delivering nucleic acid to cells for gene therapy and antisense
PT treatment.
XX Claim 16; Page 151; 293pp; English.
XX AAW96827-77 represent nucleic acid binding domains derived from human
CC apolipoprotein B-100 (apob-100). Apob-100 is a major apoprotein component
CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
CC sequence can be used in the composition of the invention. The
CC specification describes a composition that comprises LDL and
CC apolipoproteins for the binding and in vivo transport of nucleic acids.
CC The composition is used to deliver nucleic acids to eukaryotic cells, in
CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
CC molecule (or ribozyme). Specifically they are used for gene therapy of
CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
CC fibrosis and arteriosclerosis
XX SQ Sequence 51 AA;
Query Match 89.8%; Score 44; DB 2; Length 51;
Best Local Similarity 90.0%; Pred. No. 0.46;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TELTRKRGK 10
Db 6 TRLTRKRGK 15
RESULT 14
ABB37687
ID ABB37687 standard; peptide; 343 AA.
XX ABB37687;
AC ABB37687;
XX 04-FEB-2002 (first entry)
XX Peptide #5193 encoded by human foetal liver single exon probe.
XX Human; foetal liver; gene expression; single exon nucleic acid probe.
XX Homo sapiens.
XX

PN WO200157277-A2.
XX 09-AUG-2001.
XX 30-JAN-2001; 2001WO-US000669.
XX 04-FEB-2000; 2000US-0180312P.
XX 26-MAY-2000; 2000US-0207456P.
XX 30-JUN-2000; 2000US-00608408.
XX 03-AUG-2000; 2000US-00632366.
XX 21-SEP-2000; 2000US-0234687P.
XX 27-SEP-2000; 2000US-0236359P.
XX 04-OCT-2000; 2000GB-00024263.
XX (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-483447/52.
XX Human genome-derived single exon nucleic acid probes useful for analyzing
PT gene expression in human fetal liver.
XX Claim 27; SEQ ID NO 30322; 639pp + Sequence Listing; English.
XX The invention relates to a single exon nucleic acid probe for measuring
CC human gene expression in a sample derived from human foetal liver. The
CC single exon nucleic acid probes may be used for predicting, measuring and
CC displaying gene expression in samples derived from human fetal liver. The
CC present sequence is a peptide encoded by a single exon nucleic acid probe
CC of the invention. Note: The sequence data for this patent did not form
CC part of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX SQ Sequence 343 AA;
Query Match 89.8%; Score 44; DB 4; Length 343;
Best Local Similarity 90.0%; Pred. No. 3.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TELTRKRGK 10
Db 169 TRLTRKRGK 178
RESULT 15
ABG52504
ID ABG52504 standard; peptide; 343 AA.
XX ABG52504;
XX 25-FEB-2003 (first entry)
XX Human liver peptide, SEQ ID No 31152.
XX Human; liver; cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
XX hypercholesterolaemia; coronary heart disease.
XX Homo sapiens.
XX WO200157273-A2.
XX 09-AUG-2001.
XX 30-JAN-2001; 2001WO-US000664.
XX 04-FEB-2000; 2000US-0180312P.
XX 26-MAY-2000; 2000US-0207456P.
XX 30-JUN-2000; 2000US-00608408.
XX 03-AUG-2000; 2000US-00632366.
XX 21-SEP-2000; 2000US-0234687P.
XX 27-SEP-2000; 2000US-0236359P.
XX 04-OCT-2000; 2000GB-00024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.
 XX PA Penn SG, Hanzel DK, Chen W, Rank DR;
 XX PI WPI; 2001-48898/53.
 XX DR Human genome-derived single exon nucleic acid probes useful for analyzing
 XX PT gene expression in human adult liver.
 XX PT
 XX PS Claim 27; SEQ ID NO 31152; 658pp; English.
 XX CC The invention relates to a single exon nucleic acid probe (SENP) (I) for
 CC measuring human gene expression in a sample derived from human adult
 CC liver, comprising one of 13109 defined nucleotide sequences given in the
 CC specification (or complements/ fragments). The probe hybridises at high
 CC stringency to a nucleic acid molecule expressed in the human adult liver.
 CC (i) may be used for predicting, measuring and displaying gene expression
 CC in samples derived from human adult liver. The genes identified may be
 CC involved in genetic liver diseases such as cirrhosis
 CC hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
 CC associated with coronary heart disease. ABG47348-ABG5930 represent human
 CC liver single exon encoded peptides of the invention. Note: The sequence
 CC information for this patent does not appear in the printed specification
 CC but was obtained in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX SQ Sequence 343 AA;

Query Match 89.8%; Score 44; DB 4; Length 343;
 Best Local Similarity 90.0%; Pred. No. 3.1;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TELTRKRLK 10
 Db 169 TELTRKRLK 178
 | |||||
 | |||||

Search completed: January 13, 2005, 01:43:01
 Job time : 78.3798 secs

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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:43:16 ; Search time 10.6557 Seconds
(without alignments)
47.947 Million cell updates/sec

Title: US-09-823-418-11

Perfect score: 49

Sequence: 1 TELTRKRLK 10

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 219722 seqs, 51091598 residues

Total number of hits satisfying chosen parameters: 219722

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Pending Patents_AA New:*

- 1: /cgn2_6/ptodata/1/paa/PCT_NEW_COMB.pep:*
- 2: /cgn2_6/ptodata/1/paa/US06_NEW_COMB.pep:*
- 3: /cgn2_6/ptodata/1/paa/US07_NEW_COMB.pep:*
- 4: /cgn2_6/ptodata/1/paa/US08_NEW_COMB.pep:*
- 5: /cgn2_6/ptodata/1/paa/US09_NEW_COMB.pep:*
- 6: /cgn2_6/ptodata/1/paa/US10_NEW_COMB.pep:*
- 7: /cgn2_6/ptodata/1/paa/US11_NEW_COMB.pep:*
- 8: /cgn2_6/ptodata/1/paa/US60_NEW_COMB.pep:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------|-----------------------|
| 1 | 44 | 89.8 | 42 | 6 | US-10-861-779-1 |
| 2 | 44 | 89.8 | 3000 | 8 | US-60-636-722-924 |
| 3 | 44 | 89.8 | 3262 | 8 | US-60-636-722-923 |
| 4 | 44 | 89.8 | 4560 | 6 | US-10-398-200-2 |
| 5 | 44 | 89.8 | 4563 | 6 | US-10-868-577A-25 |
| 6 | 44 | 89.8 | 4563 | 8 | US-60-636-722-919 |
| 7 | 44 | 89.8 | 4563 | 8 | US-60-636-722-921 |
| 8 | 44 | 89.8 | 4563 | 8 | US-60-636-722-922 |
| 9 | 44 | 89.8 | 4563 | 8 | US-60-636-722-925 |
| 10 | 41 | 83.7 | 720 | 6 | US-10-990-328-11685 |
| 11 | 41 | 83.7 | 720 | 6 | US-10-990-328-11686 |
| 12 | 41 | 83.7 | 720 | 6 | US-10-990-328-11687 |
| 13 | 41 | 83.7 | 720 | 6 | US-10-990-328-11688 |
| 14 | 41 | 83.7 | 720 | 8 | US-60-636-720-278 |
| 15 | 41 | 83.7 | 720 | 8 | US-60-636-720-279 |
| 16 | 41 | 83.7 | 720 | 8 | US-60-636-720-280 |
| 17 | 41 | 83.7 | 720 | 8 | US-60-636-720-281 |
| 18 | 41 | 83.7 | 720 | 8 | US-60-636-720-282 |
| 19 | 41 | 83.7 | 720 | 8 | US-60-636-720-283 |
| 20 | 34 | 69.4 | 1040 | 6 | US-10-777-288A-2245 |
| 21 | 34 | 69.4 | 1284 | 6 | US-10-408-765-1008 |
| 22 | 33 | 67.3 | 205 | 6 | US-10-732-923-16259 |
| 23 | 33 | 67.3 | 227 | 6 | US-10-732-923-16246 |
| 24 | 33 | 67.3 | 297 | 1 | PCT-US02-09107B-65100 |
| 25 | 33 | 67.3 | 548 | 1 | PCT-US02-09107B-65704 |

| | | | | | | |
|----|----|------|------|---|---------------------|-------------------|
| 26 | 33 | 67.3 | 697 | 6 | US-10-999-233-18 | Sequence 18, Appl |
| 27 | 33 | 67.3 | 745 | 6 | US-10-999-233-16 | Sequence 16, Appl |
| 28 | 33 | 67.3 | 791 | 6 | US-10-999-233-22 | Sequence 22, Appl |
| 29 | 33 | 67.3 | 839 | 6 | US-10-999-233-20 | Sequence 20, Appl |
| 30 | 33 | 67.3 | 886 | 6 | US-10-732-923-3311 | Sequence 3311, Ap |
| 31 | 33 | 67.3 | 963 | 6 | US-10-408-765-2504 | Sequence 2504, Ap |
| 32 | 33 | 67.3 | 966 | 6 | US-10-408-765-2155 | Sequence 2155, Ap |
| 33 | 33 | 67.3 | 1175 | 6 | US-10-999-233-26 | Sequence 26, Appl |
| 34 | 33 | 67.3 | 1175 | 6 | US-10-999-233-30 | Sequence 30, Appl |
| 35 | 33 | 67.3 | 1259 | 6 | US-10-999-233-4 | Sequence 4, Appli |
| 36 | 33 | 67.3 | 1298 | 6 | US-10-999-233-24 | Sequence 24, Appl |
| 37 | 33 | 67.3 | 1298 | 6 | US-10-999-233-28 | Sequence 28, Appl |
| 38 | 33 | 67.3 | 1307 | 6 | US-10-999-233-2 | Sequence 2, Appli |
| 39 | 32 | 65.3 | 314 | 6 | US-10-990-328-10596 | Sequence 10596, A |
| 40 | 32 | 65.3 | 414 | 6 | US-10-990-328-10595 | Sequence 10595, A |
| 41 | 32 | 65.3 | 414 | 6 | US-10-990-328-10597 | Sequence 10597, A |
| 42 | 32 | 65.3 | 414 | 8 | US-60-631-993-330 | Sequence 330, App |
| 43 | 32 | 65.3 | 1380 | 6 | US-10-952-698-164 | Sequence 164, App |
| 44 | 32 | 65.3 | 1388 | 1 | PCT-US04-39788-2 | Sequence 2, Appli |
| 45 | 32 | 65.3 | 1975 | 1 | PCT-US04-37204-4754 | Sequence 4754, Ap |

ALIGNMENTS

RESULT 1
US-10-861-779-1
; Sequence 1, Application US/10861779
; GENERAL INFORMATION:
; APPLICANT: Verma, Inder M.
; TITLE OF INVENTION: Compositions and Methods For Targeting a
; TITLE OF INVENTION: Polypeptide To the Central Nervous System
; FILE REFERENCE: 66671-131
; CURRENT APPLICATION NUMBER: US/10/861,779
; CURRENT FILING DATE: 2004-06-04
; PRIOR APPLICATION NUMBER: 60/476,482
; PRIOR FILING DATE: 2003-06-05
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 1
; LENGTH: 42
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-861-779-1

Query Match 89.8%; Score 44; DB 6; Length 42;
Best Local Similarity 90.0%; Pred. No. 0.066;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLK 10
| | | | | | | | | |
Db 16 TELTRKRLK 25

RESULT 2
US-60-636-722-924
; Sequence 924, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 924
; LENGTH: 3000
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-924

Query Match 89.8%; Score 44; DB 8; Length 3000;

Best Local Similarity 90.0%; Pred. No. 5.3;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 2337 TRLTRKRGK 2346

RESULT 3

US-60-636-722-923
; Sequence 923, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 923
; LENGTH: 3262
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-923

Query Match 89.8%; Score 44; DB 8; Length 3262;
Best Local Similarity 90.0%; Pred. No. 5.8;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 2084 TRLTRKRGK 2093

RESULT 4

US-10-398-200-2
; Sequence 2, Application US/10398200
; GENERAL INFORMATION:
; APPLICANT: AGNELLO, VINCENT
; TITLE OF INVENTION: METHOD OF INHIBITING INFECTION BY HCV, OTHER
; TITLE OF INVENTION: FLAVIVIRIDAE VIRUSES, AND ANY OTHER VIRUS THAT
; TITLE OF INVENTION: COMPLEXES TO LOW DENSITY LIPOPROTEIN OR TO VERY LOW
; TITLE OF INVENTION: DENSITY LIPOPROTEIN IN BLOOD BY PREVENTING VIRAL ENTRY
; TITLE OF INVENTION: INTO A CELL
; FILE REFERENCE: 1513-PCT-00
; CURRENT APPLICATION NUMBER: US/10/398,200
; CURRENT FILING DATE: 2003-04-03
; PRIOR APPLICATION NUMBER: 60/243,594
; PRIOR FILING DATE: 2000-10-25
; NUMBER OF SEQ ID NOS: 3
; SOFTWARE: PatentIn Ver. 2.1
; SEQ ID NO 2
; LENGTH: 4560
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-398-200-2

Query Match 89.8%; Score 44; DB 6; Length 4560;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 3382 TRLTRKRGK 3391

RESULT 5

US-10-868-577A-25
; Sequence 25, Application US/10868577A
; GENERAL INFORMATION:
; APPLICANT: Alitalo et al.
; TITLE OF INVENTION: HEPARIN BINDING VEGFR-3 LIGANDS
; FILE REFERENCE: 28967/39359A

; CURRENT APPLICATION NUMBER: US/10/868,577A
; CURRENT FILING DATE: 2004-06-14
; PRIOR APPLICATION NUMBER: US 60/478,390
; PRIOR FILING DATE: 2003-06-12
; PRIOR APPLICATION NUMBER: US 10/669,176
; PRIOR FILING DATE: 2003-09-23
; NUMBER OF SEQ ID NOS: 69
; SOFTWARE: PatentIn version 3.2
; SEQ ID NO 25
; LENGTH: 4563
; TYPE: PRT
; ORGANISM: Homo sapiens
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (32)-(126)
; OTHER INFORMATION: heparin binding domain
; FEATURE:
; NAME/KEY: misc feature
; LOCATION: (3161)..(3236)
; OTHER INFORMATION: heparin binding domain
US-10-868-577A-25

Query Match 89.8%; Score 44; DB 6; Length 4563;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 3385 TRLTRKRGK 3394

RESULT 6

US-60-636-722-919
; Sequence 919, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 919
; LENGTH: 4563
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-919

Query Match 89.8%; Score 44; DB 8; Length 4563;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 3385 TRLTRKRGK 3394

RESULT 7

US-60-636-722-921
; Sequence 921, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 921
; LENGTH: 4563
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-921

```
Query Match      89.8%; Score 44; DB 8; Length 4563;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 3385 TELTRKGLK 3394

RESULT 8
US-60-636-722-922
; Sequence 922, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 922
; LENGTH: 4563
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-922

Query Match      89.8%; Score 44; DB 8; Length 4563;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 3385 TELTRKGLK 3394

RESULT 9
US-60-636-722-925
; Sequence 925, Application US/60636722
; GENERAL INFORMATION:
; APPLICANT: JOSELOFF, Elizabeth et al.
; TITLE OF INVENTION: LUNG DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001568
; CURRENT APPLICATION NUMBER: US/60/636,722
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 2719
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 925
; LENGTH: 4563
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-722-925

Query Match      89.8%; Score 44; DB 8; Length 4563;
Best Local Similarity 90.0%; Pred. No. 8.1;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 3385 TELTRKGLK 3394

RESULT 10
US-10-990-328-11685
; Sequence 11685, Application US/10990328
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele
; TITLE OF INVENTION: POLYMORPHISMS IN NUCLEIC ACID MOLECULES
; TITLE OF INVENTION: ENCODING HUMAN ENZYME PROTEINS, METHODS OF DETECTION AND
; FILE REFERENCE: CL001495
; CURRENT APPLICATION NUMBER: US/10/990,328
; CURRENT FILING DATE: 2004-11-17
```

```
; NUMBER OF SEQ ID NOS: 558824
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11685
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-990-328-11685

Query Match      83.7%; Score 41; DB 6; Length 720;
Best Local Similarity 80.0%; Pred. No. 4.7;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 633 TELARKKGLK 642

RESULT 11
US-10-990-328-11686
; Sequence 11686, Application US/10990328
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele
; TITLE OF INVENTION: POLYMORPHISMS IN NUCLEIC ACID MOLECULES
; TITLE OF INVENTION: ENCODING HUMAN ENZYME PROTEINS, METHODS OF DETECTION AND
; FILE REFERENCE: CL001495
; CURRENT APPLICATION NUMBER: US/10/990,328
; CURRENT FILING DATE: 2004-11-17
; NUMBER OF SEQ ID NOS: 558824
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11686
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-990-328-11686

Query Match      83.7%; Score 41; DB 6; Length 720;
Best Local Similarity 80.0%; Pred. No. 4.7;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 633 TELARKKGLK 642

RESULT 12
US-10-990-328-11687
; Sequence 11687, Application US/10990328
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele
; TITLE OF INVENTION: POLYMORPHISMS IN NUCLEIC ACID MOLECULES
; TITLE OF INVENTION: ENCODING HUMAN ENZYME PROTEINS, METHODS OF DETECTION AND
; FILE REFERENCE: CL001495
; CURRENT APPLICATION NUMBER: US/10/990,328
; CURRENT FILING DATE: 2004-11-17
; NUMBER OF SEQ ID NOS: 558824
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 11687
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-990-328-11687

Query Match      83.7%; Score 41; DB 6; Length 720;
Best Local Similarity 80.0%; Pred. No. 4.7;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKGLK 10
Db 633 TELARKKGLK 642
```

RESULT 13

US-10-990-328-11688
; Sequence 11688, Application US/10990328
; GENERAL INFORMATION:
; APPLICANT: CARGILL, Michele
; TITLE OF INVENTION: POLYMORPHISMS IN NUCLEIC ACID MOLECULES
; TITLE OF INVENTION: ENCODING HUMAN ENZYME PROTEINS, METHODS OF DETECTION AND
; FILE REFERENCE: CL001495
; CURRENT APPLICATION NUMBER: US/10/990,328
; CURRENT FILING DATE: 2004-11-17
; NUMBER OF SEQ ID NOS: 558824
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 11688
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-990-328-11688

Query Match 83.7%; Score 41; DB 6; Length 720;

Best Local Similarity 80.0%; Pred. No. 4.7;

Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10
||| ||:|
Db 633 TELARKKGLK 642

RESULT 14

US-60-636-720-278
; Sequence 278, Application US/60636720
; GENERAL INFORMATION:
; APPLICANT: DOMON, Bruno et al.
; TITLE OF INVENTION: COLON DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001567
; CURRENT APPLICATION NUMBER: US/60/636,720
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 4325
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 278
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-720-278

Query Match 83.7%; Score 41; DB 8; Length 720;

Best Local Similarity 80.0%; Pred. No. 4.7;

Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10
||| ||:|
Db 633 TELARKKGLK 642

RESULT 15

US-60-636-720-279
; Sequence 279, Application US/60636720
; GENERAL INFORMATION:
; APPLICANT: DOMON, Bruno et al.
; TITLE OF INVENTION: COLON DISEASE TARGETS AND USES THEREOF
; FILE REFERENCE: CL001567
; CURRENT APPLICATION NUMBER: US/60/636,720
; CURRENT FILING DATE: 2004-12-17
; NUMBER OF SEQ ID NOS: 4325
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 279
; LENGTH: 720
; TYPE: PRT
; ORGANISM: Homo sapiens
US-60-636-720-279

Query Match 83.7%; Score 41; DB 8; Length 720;

Best Local Similarity 80.0%; Pred. No. 4.7;

Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10

||| ||:|
Db 633 TELARKKGLK 642

Search completed: January 13, 2005, 02:20:03
Job time : 10.6557 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:30:05 ; Search time 14.4262 Seconds
(without alignments)
66.696 Million cell updates/sec

Title: US-09-823-418-11
Perfect score: 49
Sequence: 1 TELTRKRLGLK 10
Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues
Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79.*
1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|----------|--------------------|
| 1 | 44 | 89.8 | 275 | 2 E60950 | apolipoprotein B-1 |
| 2 | 44 | 89.8 | 596 | 2 S32802 | apolipoprotein B - |
| 3 | 44 | 89.8 | 4563 | 1 LPHUB | apolipoprotein B-1 |
| 4 | 40 | 81.6 | 269 | 2 C60950 | apolipoprotein B-1 |
| 5 | 40 | 81.6 | 274 | 2 A60950 | apolipoprotein B-1 |
| 6 | 40 | 81.6 | 779 | 2 JH0102 | apolipoprotein B - |
| 7 | 37 | 75.5 | 309 | 1 E65112 | hypothetical 34.6 |
| 8 | 37 | 75.5 | 309 | 2 E85985 | hypothetical prote |
| 9 | 37 | 75.5 | 309 | 2 B91140 | hypothetical prote |
| 10 | 36 | 73.5 | 254 | 2 H95070 | hypothetical prote |
| 11 | 36 | 73.5 | 264 | 2 F97938 | hypothetical prote |
| 12 | 35 | 71.4 | 149 | 2 E72338 | conserved hypothet |
| 13 | 35 | 71.4 | 214 | 2 A82934 | transcription regu |
| 14 | 35 | 71.4 | 219 | 2 B98348 | probable transcrip |
| 15 | 35 | 71.4 | 1112 | 2 T47784 | hypothetical prote |
| 16 | 34 | 69.4 | 550 | 1 WZCKI | isocitrate lyase (|
| 17 | 34 | 69.4 | 784 | 2 JH0101 | apolipoprotein B-1 |
| 18 | 34 | 69.4 | 1058 | 2 S65460 | apolipoprotein B - |
| 19 | 34 | 69.4 | 1429 | 2 T41699 | C3-domain family p |
| 20 | 34 | 69.4 | 1778 | 2 JH0382 | apolipoprotein B - |
| 21 | 34 | 69.4 | 2629 | 2 I46569 | apolipoprotein B - |
| 22 | 33 | 67.3 | 219 | 2 A53305 | pentose-5-phosphat |
| 23 | 33 | 67.3 | 227 | 2 H84614 | probable MAD3-box |
| 24 | 33 | 67.3 | 235 | 2 E86826 | amino acid permeas |
| 25 | 33 | 67.3 | 240 | 2 A70463 | rRNA methylase - A |
| 26 | 33 | 67.3 | 249 | 2 T16924 | hypothetical prote |
| 27 | 33 | 67.3 | 309 | 2 AH0906 | conserved hypothet |
| 28 | 33 | 67.3 | 443 | 2 D72383 | NADH oxidase - The |
| 29 | 33 | 67.3 | 548 | 2 G81959 | conserved hypothet |

30 33 67.3 583 2 T04531 nine-cis-epoxycaro
31 33 67.3 642 2 T05683 hypothetical prote
32 33 67.3 963 2 T26032 hypothetical prote
33 33 67.3 1091 2 T35822 probable regulator
34 33 67.3 1253 1 A44400 myosin heavy chain
35 33 67.3 1254 2 A54818 myosin-VI [similar
36 33 67.3 1265 2 A59299 unconventional myo
37 33 67.3 1407 2 S59823 probable membrane
38 33 67.3 1615 2 JC6510 ras-responsive ele
39 33 67.3 2044 2 T13704 still life protein
40 33 67.3 2064 2 T13707 still life protein
41 32 65.3 107 2 S12607 salivary glue prot
42 32 65.3 112 2 S33822 salivary glue prot
43 32 65.3 172 2 F69506 probable 2-oxoisov
44 32 65.3 183 2 E84119 ATP synthase delta
45 32 65.3 237 2 T19702 hypothetical prote

ALIGNMENTS

RESULT 1

E60950
apolipoprotein B-100 - chicken (fragment)
C/Species: Gallus gallus (chicken)
C/Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C/Accession: E60950
R/Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A/Title: A cross-species comparison of the apolipoprotein B domain that binds to the LD
A/Reference number: A60950; MUID:90324804; PMID:2373961
A/Accession: E60950
A/Molecule type: mRNA
A/Residues: 1-275 <LAW>
A/Cross-references: UNIPROT:Q7LZ77
A/Superfamily: apolipoprotein B
C/Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 89.8%; Score 44; DB 2; Length 275;
Best Local Similarity 90.0%; Pred. No. 0.49;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLGLK 10
| | | | | | | | | |
Db 221 TSLTRKRLGLK 230

RESULT 2

S32802
apolipoprotein B - crab-eating macaque (fragment)
C/Species: Macaca fascicularis (crab-eating macaque)
C/Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Jul-2004
C/Accession: S32802
R/Pape, M.B.; Castle, C.K.; Murray, R.W.; Funk, G.M.; Hunt, C.E.; Marotti, K.R.; Melchior
Biochim. Biophys. Acta 1086, 326-334, 1991
A/Title: Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional r
A/Reference number: S32802; MUID:92075708; PMID:1742325
A/Accession: S32802
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-596 <PAP>
A/Cross-references: UNIPROT:Q28473; EMBL:X15737; NID:G38047; PIDN:CAA33755.1; PID:G9301
C/Superfamily: apolipoprotein B

Query Match 89.8%; Score 44; DB 2; Length 596;
Best Local Similarity 90.0%; Pred. No. 0.98;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLGLK 10
| | | | | | | | | |
Db 226 TRLTRKRLGLK 235

RESULT 3

LPHUB

apolipoprotein B-100 precursor - human
 N1:Contains: apolipoprotein B-26; apolipoprotein B-48; apolipoprotein B-74
 C:Species: Homo sapiens (man)
 C1:Date: 28-Dec-1987 #sequence revision 28-Dec-1987 #text_change 09-Jul-2004
 C1:Accession: A27850; A25267; A25263; A24320; A24684; A23817; A25774; A264452; I61909; I59510; I39474; I39469; I84624; I37179; PS0058
 R1: Ludwig, E.H.; Blackhart, B.D.; Pierotti, V.R.; Caiati, L.; Fortier, C.; Knott, T.; SocDNA 6, 363-372, 1987
 A1:Title: DNA sequence of the human apolipoprotein B gene.
 A1:Reference number: A27850; MUID:88003974; PMID:3652307
 A1:Accession: A27850
 A1:Molecule type: DNA
 A1:Residues: 1-617, 'A', 619-1929, 'F', 1931-3318, 'D', 3320-3426, 'T', 3428-3431, 'Q', 3433-3731, 'R'; Knott, T.J.; Wallis, S.C.; Powell, L.M.; Pease, R.J.; Lusis, A.J.; Blackhart, B.; McGa
 Nucleic Acids Res. 14, 7501-7503, 1986
 R1:Cladara, C.; Hadzopoulou-Cladara, M.; Nolte, R.T.; Atkinson, D.; Zannis, V.I.
 EMBO J. 5, 3495-3507, 1986
 A1:Title: The complete sequence and structural analysis of human apolipoprotein B-100: re
 A1:Reference number: A91058; MUID:87161758; PMID:3030729
 A1:Accession: A25679
 A1:Molecule type: mRNA
 A1:Residues: 1-11, 15-2339, 'S', 2541-3823, 'R', 3825-4563 <CLA>
 A1:Note: 1109-Asp was also found
 R1:Knott, T.J.; Wallis, S.C.; Powell, L.M.; Pease, R.J.; Lusis, A.J.; Blackhart, B.; McGa
 Nucleic Acids Res. 14, 7501-7503, 1986
 A1:Title: Complete cDNA and derived protein sequence of human apolipoprotein B-100.
 A1:Reference number: A93639; MUID:87016385; PMID:3763409
 A1:Accession: A25263
 A1:Molecule type: mRNA
 A1:Residues: 1-272, 'N', 274-617, 'A', 619-1217, 'E', 1219-2091, 'V', 2093-2364, 'T', 2366-2679, 'Q'
 A1:Cross-references: GB:X04506; NID:G34330; PIDN:CAA28191.1; PID:G34331
 R1:Law, S.W.; Grant, S.M.; Higuchi, K.; Hospattankar, A.; Lackner, K.; Lee, N.; Brewer J
 Proc. Natl. Acad. Sci. U.S.A. 83, 8143-8146, 1986
 A1:Title: Human liver apolipoprotein B-100 cDNA: complete nucleic acid and derived amino
 A1:Reference number: A94134; MUID:87041416; PMID:3464946
 A1:Accession: A25267
 A1:Molecule type: mRNA
 A1:Residues: 1-617, 'A', 619-703, 'P', 705-792, 'R', 794-1270, 'S', 1272-1866, 'G', 1868-2036, 'N', 2
 A1:Note: the codons given for residues 704, 793, 1271, 2037, 2933, 3286, 3782, 4188, and
 R1:Chen, S.H.; Yang, C.Y.; Chen, P.F.; Setzer, D.; Tanimura, M.; Li, W.H.; Gotto Jr., A.M
 J. Biol. Chem. 261, 12918-12921, 1986
 A1:Title: The complete cDNA and amino acid sequence of human apolipoprotein B-100.
 A1:Reference number: A92556; MUID:87008488; PMID:3759943
 A1:Accession: A25266
 A1:Molecule type: mRNA
 A1:Residues: 1-97, 'I', 99-328, 'V', 330-644, 'I', 646-918, 'P', 920-3318, 'D', 3320-3426, 'T', 3428-
 9-4132, 'G', 4134-4180, 'E', 4182-4563 <CHE>
 A1:Cross-references: GB:J02610; NID:G178803; PIDN:AAA35549.1; PID:G178804
 A1:Note: a total of 2366 residues were confirmed by direct sequencing of tryptic peptides
 R1:Protter, A.A.; Hardman, D.A.; Sato, K.Y.; Schilling, J.W.; Yamanaka, M.; Hort, Y.J.; H
 Proc. Natl. Acad. Sci. U.S.A. 83, 5678-5682, 1986
 A1:Title: Analysis of cDNA clones encoding the entire B-26 region of human apolipoprotein
 A1:Reference number: A24320; MUID:86287319; PMID:3461454
 A1:Accession: A24320
 A1:Molecule type: mRNA
 A1:Residues: 1-97, 'I', 99-617, 'A', 619-941, 'Y', 941-1138, 'P', 1138-1138, 'PTGRLNCFNSGLICSLWLSHSPQ
 A1:Cross-references: GB:M14081; NID:G178795; PIDN:AAA51752.1; PID:G553189
 R1:Law, S.W.; Lackner, K.J.; Hospattankar, A.V.; Anchors, J.M.; Sakaguchi, A.Y.; Naylor,
 Proc. Natl. Acad. Sci. U.S.A. 82, 8340-8344, 1985
 A1:Title: Human apolipoprotein B-100: cloning, analysis of liver mRNA, and assignment of
 A1:Reference number: A24684; MUID:86094221; PMID:3001697
 A1:Accession: A24684
 A1:Molecule type: mRNA
 A1:Residues: 485-617, 'A', 619-1044 <LA2>
 A1:Cross-references: GB:M12480; NID:G178791; PIDN:AAA51751.1; PID:G178792
 R1:Protter, A.A.; Hardman, D.A.; Schilling, J.W.; Miller, J.; Appleby, V.; Chen, G.C.; Ki
 Proc. Natl. Acad. Sci. U.S.A. 83, 1467-1471, 1986
 A1:Title: Isolation of a cDNA clone encoding the amino-terminal region of human apolipop
 A1:Reference number: A94088; MUID:86149325; PMID:3513177
 A1:Accession: A23817
 A1:Molecule type: mRNA

A1:Residues: 1-291 <PRO>
 A1:Cross-references: GB:M12681; NID:G178797; PIDN:AAA51753.1; PID:G178798
 R1:Deeb, S.S.; Motulsky, A.G.; Albers, J.J.
 Proc. Natl. Acad. Sci. U.S.A. 82, 4983-4986, 1985
 A1:Title: A partial cDNA clone for human apolipoprotein B.
 A1:Reference number: A25774; MUID:85270450; PMID:3860836
 A1:Accession: A25774
 A1:Molecule type: mRNA
 A1:Residues: 709-791, 'SSSWKAASHGCPHSGD', 810-906 <DEE>
 A1:Cross-references: GB:K03175; NID:G178821; PIDN:AAA51759.1; PID:G178822
 R1:Carlsson, P.; Darnfors, C.; Olofsson, S.O.; Bjursell, G.
 Gene 49, 29-51, 1986
 A1:Title: Analysis of the human apolipoprotein B gene; complete structure of the B-74 re
 A1:Reference number: A91565; MUID:87191999; PMID:2883086
 A1:Accession: A26533
 A1:Molecule type: mRNA
 A1:Residues: 1282-2721, 2742-3290, 'L', 3292-3336, 'N', 3338-3948, 'F', 3950-3963, 'Y', 3965-4180, 'R';
 A1:Cross-references: GB:M15421; NID:G178817; PIDN:AAA51758.1; PID:G178818
 R1:Hardman, D.A.; Protter, A.A.; Chen, G.C.; Schilling, J.W.; Sato, K.Y.; Lau, K.; Yaman
 Biochemistry 26, 5478-5486, 1987
 A1:Title: Structural comparison of human apolipoproteins B-48 and B-100.
 A1:Reference number: A29671; MUID:88050832; PMID:3676265
 A1:Accession: A29671
 A1:Molecule type: mRNA
 A1:Residues: 1671-2323, 'PYW', 2327-2352, 'H', 2354-2398 <HAR>
 A1:Cross-references: GB:M17367; NID:G178731; PIDN:AAA51741.1; PID:G178732
 R1:Shoulders, C.C.; Myant, N.B.; Sidoli, A.; Rodriguez, J.C.; Cortese, C.; Baralle, F.E.;
 Atherosclerosis 58, 277-289, 1985
 A1:Title: Molecular cloning of human LDL apolipoprotein B cDNA. Evidence for more than o
 A1:Reference number: A90084; MUID:86130855; PMID:3841481
 A1:Accession: A23287
 A1:Molecule type: mRNA
 A1:Residues: 3846-4298 <SHO>
 R1:Pfitzner, R.; Wagener, R.; Stoffel, W.
 Biol. Chem. Hoppe-Seyler 367, 1077-1083, 1986
 A1:Title: Isolation, expression and characterization of a human apolipoprotein B 100-spe
 A1:Reference number: A25572; MUID:87076044; PMID:3024665
 A1:Accession: A25572
 A1:Molecule type: mRNA
 A1:Residues: 4219-4337, 'S', 4339-4563 <PFI>
 A1:Cross-references: GB:M36676
 R1:Wei, C.F.; Chen, S.H.; Yang, C.Y.; Marcel, Y.L.; Milne, R.W.; Li, W.H.; Sparrow, J.T.
 Proc. Natl. Acad. Sci. U.S.A. 82, 7265-7269, 1985
 A1:Reference number: A24738; MUID:86042646; PMID:2932736
 A1:Accession: A24738
 A1:Molecule type: mRNA
 A1:Residues: 'N', 3729-3731, 'I', 3733-3875, 'A', 3877-3948, 'F', 3950-3963, 'Y', 3965-3982, 'S', 3
 A1:Cross-references: GB:M12413; NID:G178735; PIDN:AAA51742.1; PID:G178736
 R1:Chen, S.H.; Habib, G.; Yang, C.Y.; Gu, Z.W.; Lee, B.R.; Weng, S.; Silberman, S.R.; Ca
 Science 238, 363-366, 1987
 A1:Title: Apolipoprotein B-48 is the product of a messenger RNA with an organ-specific i
 A1:Reference number: A40133; MUID:88018019; PMID:3659919
 A1:Accession: B40133
 A1:Molecule type: mRNA
 A1:Residues: 2165-2179 <CH1>
 A1:Cross-references: GB:M18036; NID:G178799; PIDN:AAA51754.1; PID:G178800
 A1:Note: this mRNA includes the stop codon of the organ-specific mRNA for apo48
 A1:Accession: A40133
 A1:Molecule type: protein
 A1:Residues: 51-75, 101-110, 129-139, 158-174, 197-207, 276-287, 298-304, 306-314, 526-532, 538-5
 36, 1486-1498, 1537-1556, 1563-1572, 1601-1610, 1647-1661, 1697-1724, 1770-1781, 1859-1897, 1968
 A1:Note: these fragments were derived from apo48
 R1:Hardman, D.A.; Protter, A.A.; Schilling, J.W.; Kane, J.P.
 Biochem. Biophys. Res. Commun. 149, 1214-1219, 1987
 A1:Title: Carboxyl terminal analysis of human B-48 protein confirms the novel mechanism i
 A1:Reference number: A28002; MUID:88106542; PMID:3426612
 A1:Accession: A28002
 A1:Molecule type: mRNA
 A1:Residues: 2129-2179, 2181-2235 <HA2>
 A1:Cross-references: GB:M18471
 A1:Experimental source: intestine
 A1:Note: this mRNA from intestine includes a stop codon created by RNA editing in place
 R1:Mehrabian, M.; Schumaker, V.N.; Fareed, G.C.; West, R.; Johnson, D.F.; Kirchgessner, J.

Nucleic Acids Res. 13, 6937-6953, 1985
A;Title: Human apolipoprotein B: identification of cDNA clones and characterization of m
A;Reference number: A24269; MUID:86041888; PMID:3903660
A;Accession: A24269
A;Molecule type: mRNA
A;Residues: 3056-3159 <MEH>
R;Cross-references: GB:X03045; NID:g28783; PIDN:CAA26850.1; PID:g929609
R;Hospatankar, A.V.; Higuchi, K.; Law, S.W.; Meglin, N.; Brewer Jr., H.B.
Biochem. Biophys. Res. Commun. 148, 279-285, 1987
A;Title: Identification of a novel in-frame translational stop codon in human intestine
A;Reference number: A29659; MUID:88049670; PMID:2445342
A;Accession: A29659
A;Molecule type: mRNA
A;Residues: 2169-2179 <HOS>
A;Note: the sequence shown represents the carboxyl end of apolipoprotein B-48
A;Note: two RNA species, 14.1kb and 7.5kb in length, were isolated from the human intest
ch encodes the 250K apOB-48, CAA encoding 2180-gln is substituted from the stop codon TAA,
R;Yang, C.; Kim, T.W.; Weng, S.; Lee, B.; Yang, M.; Gotto Jr., A.M.
Proc. Natl. Acad. Sci. U.S.A. 87, 5523-5527, 1990
A;Title: Isolation and characterization of sulfhydryl and disulfide peptides of human ap
A;Reference number: A35783; MUID:90319144; PMID:2115173
A;Contents: disulfide bonds
A;Accession: A35783
A;Molecule type: protein
A;Residues: 28-41;76-97, I', 99-100;175-193;206-215;239-249;259-266;357-399;455-490;512-5
A;Note: cysteines at positions 1112, 1422, 1505, 1662, 3761, 3917, and 4217 have free su
R;LeBoeuf, R.C.; Miller, C.; Shively, J.E.; Schumaker, V.N.; Balla, M.A.; Lusis, A.J.
FEBS Lett. 170, 105-108, 1984
A;Title: Human apolipoprotein B: partial amino acid sequence.
A;Reference number: A22006; MUID:84208786; PMID:6373369
A;Accession: A22006
A;Molecule type: protein
A;Residues: 873-892, 'K', 894-896 <LE1>
A;Accession: B22006
A;Molecule type: protein
A;Residues: 3113, 'L', 3115-3130, 'R', 3132-3133, 'P', 3135-3136, 'R' <LE2>
R;Blackhart, B.D.; Ludwig, E.M.; Pierotti, V.R.; Caiati, L.; Onasch, M.A.; Wallis, S.C.;
J. Biol. Chem. 261, 15364-15367, 1986
A;Title: Structure of the human apolipoprotein B gene.
A;Reference number: A92564; MUID:8705153; PMID:2946672
A;Contents: annotation; gene structure
R;Wagener, R.; Pfitzner, R.; Stoffel, W.
Biol. Chem. Hoppe-Seyler 368, 419-425, 1987
A;Title: Studies on the organization of the human apolipoprotein B 100 gene.
A;Reference number: A90715; MUID:87271140; PMID:2886136
A;Contents: annotation; gene structure
R;Weisgraber, K.H.; Rall Jr., S.C.
J. Biol. Chem. 262, 11097-11103, 1987
A;Title: Human apolipoprotein B-100 heparin-binding sites.
A;Reference number: A92605; MUID:87280197; PMID:3301850
A;Contents: annotation; heparin binding and disulfide bond
R;Dashit, N.; Lee, D.M.; Mok, T.
Biochem. Biophys. Res. Commun. 137, 493-499, 1986
A;Title: Apolipoprotein B is a calcium binding protein.
A;Reference number: A90125; MUID:86242245; PMID:3087360
A;Contents: annotation; calcium binding
R;Carlsson, P.; Olofsson, S.O.; Bondjers, G.; Darnfors, C.; Wiklund, O.; Bjursell, G.
Nucleic Acids Res. 13, 8813-8826, 1985
A;Title: Molecular cloning of human apolipoprotein B cDNA.
A;Reference number: I37178; MUID:86093680; PMID:3841204
A;Accession: I37180

Query Match 89.8%; Score 44; DB 1; Length 4563;
Best Local Similarity 90.0%; Pred. No. 6.3;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10
| | | | | | | | | |
Db 3385 TELTRKRGK 3394

RESULT 4

C60950
apolipoprotein B-100 - golden hamster (fragment)
C;Species: Mesocricetus auratus (golden hamster)
C;Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C;Accession: C60950
R;Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A;Title: A cross-species comparison of the apolipoprotein B domain that binds to the LD
A;Reference number: A60950; MUID:90324804; PMID:2373961
A;Accession: C60950
A;Molecule type: DNA
A;Residues: 1-269 <LAW>
A;Cross-references: UNIPROT:Q60537; UNIPROT:Q60536
C;Superfamily: apolipoprotein B
C;Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 81.6%; Score 40; DB 2; Length 269;
Best Local Similarity 80.0%; Pred. No. 3;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10
: | | | | | | | | | |
Db 216 SRLTRKRGK 225

RESULT 5
A60950
apolipoprotein B-100 - rabbit (fragment)
C;Species: Oryctolagus cuniculus (domestic rabbit)
C;Date: 31-Dec-1993 #sequence_revision 09-Sep-1994 #text_change 09-Jul-2004
C;Accession: A60950
R;Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A;Title: A cross-species comparison of the apolipoprotein B domain that binds to the LD
A;Reference number: A60950; MUID:90324804; PMID:2373961
A;Accession: A60950
A;Molecule type: mRNA
A;Residues: 1-274 <LAW>
A;Cross-references: UNIPROT:Q7M2U9
A;Note: authors translated the codon GAT for residue 155 as His
C;Superfamily: apolipoprotein B
C;Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 81.6%; Score 40; DB 2; Length 274;
Best Local Similarity 80.0%; Pred. No. 3;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRGK 10
: | | | | | | | | | |
Db 221 SSLTRKRGK 230

RESULT 6
JH0102
apolipoprotein B - golden hamster (fragment)
C;Species: Mesocricetus auratus (golden hamster)
C;Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
C;Accession: JH0102
R;Smith, T.J.
submitted to GenBank, June 1990
A;Reference number: A38864
A;Accession: JH0102
A;Molecule type: DNA
A;Residues: 1-779 <SMI>
A;Cross-references: UNIPROT:Q60536; GB:M35187
A;Note: this is a revision to the sequence from reference JH0101
R;Smith, T.J.; Hautamaa, D.; Maeda, N.
Gene 87, 309-310, 1990
A;Title: Sequence of the putative low-density lipoprotein receptor-binding regions of a
A;Reference number: JH0101; MUID:90236327; PMID:2332175
A;Contents: annotation
A;Note: this sequence has been revised in reference A38864
C;Genetics:

A:Gene: apob
C:Superfamily: apolipoprotein B
C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;
F:435-445/Region: receptor binding
F:646-656/Region: receptor binding

Query Match 81.6%; Score 40; DB 2; Length 779;
Best Local Similarity 80.0%; Pred. No. 7.8;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
: |||||
Db 642 SRLTRKRGK 651

RESULT 7
E65112
Hypothetical 34.6 kD protein in arcB-gltB intergenic region - Escherichia coli (strain K12)
C:Species: Escherichia coli
C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 09-Jul-2004
C:Accession: E65112
R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; Cohen, A.; Rose, D.J.; Mau, B.; Shao, Y.
Science 277, 1453-1462, 1997
A:Title: The complete genome sequence of Escherichia coli K-12.
A:Reference number: A64720; MUID:97426617; PMID:9278503
A:Accession: E65112
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-309 <BLAT>
A:Cross-references: UNIPROT:P45476; GB:AE000400; GB:U00096; NID:G2367203; PIDN:AAC76243.
A:Experimental source: strain K-12, substrain MG1655
C:Genetics:
A:Gene: yhcC
C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 75.5%; Score 37; DB 1; Length 309;
Best Local Similarity 70.0%; Pred. No. 13;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
: |||||
Db 170 TQLARQRLK 179

RESULT 8
E85985
Hypothetical protein yhcC [imported] - Escherichia coli (strain O157:H7, substrain EDL933)
C:Species: Escherichia coli
C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 09-Jul-2004
C:Accession: E85985
R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glasner, J.D.; Rose, D.J.; Mayhew, M.W.; Miller, L.; Grotbeck, E.J.; Davis, N.W.; Lim, A.; Dimalanta, E.; Potamousis, K.; Apodaca, Nature 409, 529-533, 2001
A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
A:Reference number: A85480; MUID:21074935; PMID:11206551
A:Accession: E85985
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-309 <STO>
A:Cross-references: UNIPROT:P45476; GB:AE005174; NID:G12517832; PIDN:AAG58345.1; GSPDB:G12517832
A:Experimental source: strain O157:H7, substrain EDL933
C:Genetics:
A:Gene: yhcC
C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 75.5%; Score 37; DB 2; Length 309;
Best Local Similarity 70.0%; Pred. No. 13;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
: |||||
Db 170 TQLARQRLK 179

RESULT 9
B91140
Hypothetical protein ECs4090 [imported] - Escherichia coli (strain O157:H7, substrain R6)
C:Species: Escherichia coli
C:Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 09-Jul-2004
C:Accession: B91140
R:Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Iihil, K.; Yokoyama, K.; Han, C.G. gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H. DNA Res. 8, 11-22, 2001
A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and genome organization of the enterohemorrhagic EHEC O157:H7
A:Reference number: A99629; MUID:21156231; PMID:11256796
A:Accession: B91140
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-309 <HAY>
A:Cross-references: UNIPROT:P45476; GB:BA000007; PIDN:BA837513.1; PID:G13363563; GSPDB:BA837513.1
A:Experimental source: strain O157:H7, substrain RMD 0509952
C:Genetics:
A:Gene: ECs4090
C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 75.5%; Score 37; DB 2; Length 309;
Best Local Similarity 70.0%; Pred. No. 13;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
: |||||
Db 170 TQLARQRLK 179

RESULT 10
H95070
Hypothetical protein SP0609 [imported] - Streptococcus pneumoniae (strain TIGR4)
C:Species: Streptococcus pneumoniae
C:Date: 03-Aug-2001 #sequence_revision 03-Aug-2001 #text_change 09-Jul-2004
C:Accession: H95070
R:Tettelin, H.; Nelson, K.E.; Paulsen, I.T.; Eisen, J.A.; Read, T.D.; Peterson, S.; Helgeson, J.D.; Umayam, L.A.; White, O.; Salzberg, S.L.; Lewis, M.R.; Radune, D.; Holtzapple, M.; Hickey, E.K.; Holt, I.E. Science 293, 498-506, 2001
A:Authors: Loftus, B.J.; Yang, F.; Smith, H.O.; Venter, J.C.; Dougherty, B.A.; Morrison, A.; Title: Complete Genome Sequence of a virulent isolate of Streptococcus pneumoniae.
A:Reference number: A95000; MUID:21357209; PMID:11463916
A:Accession: H95070
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-254 <KUR>
A:Cross-references: UNIPROT:Q97S14; GB:AE005672; PIDN:AAK74761.1; PID:G14972084; GSPDB:Q97S14
A:Experimental source: strain TIGR4
C:Genetics:
A:Gene: SP0609

Query Match 73.5%; Score 36; DB 2; Length 254;
Best Local Similarity 70.0%; Pred. No. 17;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
: |||||
Db 149 TELGKKKGLK 158

RESULT 11
F97938
Hypothetical protein glnH [imported] - Streptococcus pneumoniae (strain R6)
C:Species: Streptococcus pneumoniae
C:Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 09-Jul-2004
C:Accession: F97938
R:Hoskins, J.A.; Alborn Jr., W.; Arnold, J.; Blaszcak, L.; Burgett, S.; DeHoff, B.S.; Filler, R.; LeBlanc, D.J.; Lee, L.N.; Lefkowitz, E.J.; Lu, J.; Matsushima, P.; McAhren, S.; Nisbet, P.; Sun, P.M.; Winkler, M.E. J. Bacteriol. 183, 5709-5717, 2001

A;Authors: Yang, Y.; Young-Bellido, M.; Zhao, G.; Zook, C.; Baltz, R.H.; Jaskunas, S.R.;
 A;Title: Genome of the Bacterium Streptococcus pneumoniae Strain R6.
 A;Reference number: A37872; MUID:21429245; PMID:11544234
 A;Accession: F97938
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-264 <KUR>
 A;Cross-references: UNIPROT:Q8CWT1; GB:AE007317; PIDN:AAK99338.1; PID:gl5458109; GSPDB:G
 C;Genetics:
 A;Gene: glnH

Query Match 73.5%; Score 36; DB 2; Length 264;
 Best Local Similarity 70.0%; Pred. No. 18;
 Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLGLK 10
 |||:|:|
 Db 159 TELGKKGLK 168

RESULT 12
 E72338
 conserved hypothetical protein - Thermotoga maritima (strain MSB8)
 C;Species: Thermotoga maritima
 C;Date: 11-Jun-1999 #sequence_revision 11-Jun-1999 #text_change 09-Jul-2004
 C;Accession: E72338
 R;Nelson, K.E.; Clayton, R.A.; Gilli, S.R.; Gwinn, M.L.; Dodson, R.J.; Haft, D.H.; Hickey
 Garrett, M.M.; Stewart, A.M.; Cotton, M.D.; Pratt, M.S.; Phillips, C.A.; Richardson, D.;
 C.M.

Nature 399, 323-329, 1999
 A;Title: Evidence for lateral gene transfer between Archaea and Bacteria from genome seq
 A;Reference number: A72200; MUID:99287316; PMID:10360571
 A;Accession: E72338
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-149 <ARN>
 A;Cross-references: UNIPROT:Q9WZ19; GB:AE001744; GB:AE000512; NID:g4981254; PIDN:AAD3581
 A;Experimental source: strain MSB8
 C;Genetics:
 A;Gene: TM0730
 C;Superfamily: conserved hypothetical protein HI0670

Query Match 71.4%; Score 35; DB 2; Length 149;
 Best Local Similarity 77.8%; Pred. No. 17;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 2 ELTRKRLGLK 10
 |||:|:|
 Db 111 ELRRKGLK 119

RESULT 13
 AE2934
 transcription regulator, TerR family Atu3075 [imported] - Agrobacterium tumefaciens (str
 C;Species: Agrobacterium tumefaciens
 C;Date: 11-Jan-2002 #sequence_revision 11-Jan-2002 #text_change 09-Jul-2004
 C;Accession: AE2934
 R;Wood, D.W.; Setubal, J.C.; Kaul, R.; Monks, D.; Chen, L.; Wood, G.E.; Chen, Y.; Woo, I
 erage, G.; Gillet, W.; Grant, C.; Guenther, D.; Kutayavin, T.; Levy, R.; Li, M.; McClell
 ; Karp, P.; Romero, P.; Zhang, S.
 Science 294, 2317-2323, 2001
 A;Authors: Yoo, H.; Tao, Y.; Biddle, P.; Jung, M.; Krespan, W.; Perry, M.; Gordon-Kamm,
 ster, E.W.
 A;Title: The Genome of the Natural Genetic Engineer Agrobacterium tumefaciens C58.
 A;Reference number: AB2577; MUID:21608550; PMID:11743193
 A;Accession: AE2934
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-214 <KUR>
 A;Cross-references: UNIPROT:Q8UBE1; GB:AE008689; PIDN:AAL43891.1; PID:gl7741439; GSPDB:G
 A;Experimental source: strain C58 (Dupont)
 C;Genetics:
 A;Gene: Atu3075

A;Map position: linear chromosome

Query Match 71.4%; Score 35; DB 2; Length 214;
 Best Local Similarity 87.5%; Pred. No. 23;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRG 8
 |||:|:|
 Db 2 TETTRKG 9

RESULT 14
 B98348
 probable transcription regulator PA2931 [imported] - Agrobacterium tumefaciens (strain
 C;Species: Agrobacterium tumefaciens
 C;Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 09-Jul-2004
 C;Accession: B98348
 R;Goodner, B.; Hinkle, G.; Gattung, S.; Miller, N.; Blanchard, M.; Quorollo, B.; Goldman
 A.; Liu, F.; Wollam, C.; Allinger, M.; Doughty, D.; Scott, C.; Lappas, C.; Markelz, B.
 Science 294, 2323-2328, 2001
 A;Title: Genome Sequence of the Plant Pathogen and Biotechnology Agent Agrobacterium tu
 A;Reference number: A97359; MUID:21608551; PMID:11743194
 A;Accession: B98348
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-219 <KUR>
 A;Cross-references: UNIPROT:Q8UBE1; GB:AE007870; PIDN:AAK90308.1; PID:gl5160337; GSPDB:
 C;Genetics:
 A;Gene: AGR_L_3469

A;Map position: linear chromosome

Query Match 71.4%; Score 35; DB 2; Length 219;
 Best Local Similarity 87.5%; Pred. No. 24;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRG 8
 |||:|:|
 Db 7 TETTRKG 14

RESULT 15
 T47784
 hypothetical protein F17J16.70 - Arabidopsis thaliana
 C;Species: Arabidopsis thaliana (mouse-ear cress)
 C;Date: 20-Apr-2000 #sequence_revision 20-Apr-2000 #text_change 09-Jul-2004
 C;Accession: T47784
 R;D'Angelo, M.; Vezzi, A.; Modesto, D.; Pigazzi, M.; Valle, G.; Mewes, H.W.; Rudd, S.;
 submitted to the Protein Sequence Database, April 2000
 A;Reference number: Z24476
 A;Accession: T47784
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-1112 <DAN>
 A;Cross-references: UNIPROT:Q9LYT4; EMBL:AL163527
 A;Experimental source: cultivar Columbia; BAC clone F17J16
 C;Genetics:
 A;Map position: 3
 A;Introns: 31/3; 74/1; 103/2; 153/2; 193/2; 247/3; 309/2; 367/2; 416/1; 457/2; 517/3; 5
 A;Note: F17J16.70

Query Match 71.4%; Score 35; DB 2; Length 1112;
 Best Local Similarity 87.5%; Pred. No. 11e+02;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRG 8
 |||:|:|
 Db 433 TELVRKRG 440

Search completed: January 13, 2005, 01:52:38
 Job time : 16.4262 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 78.0328 Seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-11
Perfect score: 49
Sequence: 1 TELTRKRGK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : UniProt_02.*
1: uniprot_sprot.*
2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|--------------|--------------------|
| 1 | 44 | 89.8 | 275 | 2 Q7LZ77 | Q7Lz77 gallus gall |
| 2 | 44 | 89.8 | 387 | 2 Q7YQN2 | Q7Yqn2 phalanger o |
| 3 | 44 | 89.8 | 400 | 2 Q7YQM9 | Q7Yqm9 ornithorhyn |
| 4 | 44 | 89.8 | 405 | 2 Q7YQNO | Q7Yqn0 tachyglossu |
| 5 | 44 | 89.8 | 414 | 2 Q7YQRS | Q7Yqr5 aotus vocif |
| 6 | 44 | 89.8 | 596 | 2 Q28473 | Q28473 macaca fasc |
| 7 | 44 | 89.8 | 3262 | 2 Q13788 | Q13788 homo sapien |
| 8 | 44 | 89.8 | 4563 | 1 APB HUMAN | P04114 homo sapien |
| 9 | 44 | 89.8 | 4563 | 2 Q7Z600 | Q7z600 homo sapien |
| 10 | 41 | 83.7 | 164 | 2 O6PIM8 | Q6pim8 homo sapien |
| 11 | 41 | 83.7 | 164 | 2 AAH32144 | AAh32144 homo sapi |
| 12 | 41 | 83.7 | 720 | 1 LCF3 HUMAN | O95573 homo sapien |
| 13 | 41 | 83.7 | 720 | 2 Q8IUM9 | Q8ium9 homo sapien |
| 14 | 41 | 83.7 | 720 | 2 BAB72139 | BAb72139 homo sapi |
| 15 | 40 | 81.6 | 263 | 2 Q7YQO0 | Q7Yqo0 procavia ca |
| 16 | 40 | 81.6 | 274 | 2 Q7M2U9 | Q7m2u9 oryctolagus |
| 17 | 40 | 81.6 | 304 | 2 Q7YQP9 | Q7Yqp9 echinops te |
| 18 | 40 | 81.6 | 314 | 2 Q7YQNS | Q7Yqn8 itomyx str |
| 19 | 40 | 81.6 | 316 | 2 Q7YQP3 | Q7Yqp3 nandinia bi |
| 20 | 40 | 81.6 | 318 | 2 Q7YQNS | Q7Yqn9 zalophus ca |
| 21 | 40 | 81.6 | 319 | 2 Q7YQPO | Q7Yqp0 vulpes vulp |
| 22 | 40 | 81.6 | 319 | 2 Q7YQP2 | Q7Yqp2 panthera le |
| 23 | 40 | 81.6 | 320 | 2 Q7YQP4 | Q7Yqp4 manis sp. k |
| 24 | 40 | 81.6 | 322 | 2 Q7YQPS | Q7Yqp5 manis sp. k |
| 25 | 40 | 81.6 | 339 | 2 Q7YR05 | Q7Yr05 macroscelid |
| 26 | 40 | 81.6 | 361 | 2 Q7YQP8 | Q7Yqp8 amblysomus |
| 27 | 40 | 81.6 | 364 | 2 Q7YQO1 | Q7Yqo1 dugong dugo |
| 28 | 40 | 81.6 | 386 | 2 Q7YQRI | Q7Yqr1 tupata tana |
| 29 | 40 | 81.6 | 392 | 2 Q7YR11 | Q7Yr11 tarsius syr |
| 30 | 40 | 81.6 | 411 | 2 Q7YQP7 | Q7Yqp7 ochotona pr |
| 31 | 40 | 81.6 | 421 | 2 Q7TN68 | Q7tn68 glaucomys v |

| | | | | | |
|----|----|------|-----|----------|--------------------|
| 32 | 40 | 81.6 | 422 | 2 Q7YR12 | Q7Yr12 talpa europ |
| 33 | 40 | 81.6 | 423 | 2 Q7YQO9 | Q7Yqo9 sorex monti |
| 34 | 40 | 81.6 | 426 | 2 Q7YQR2 | Q7Yqr2 alces alces |
| 35 | 40 | 81.6 | 429 | 2 Q7YQO8 | Q7Yqo8 crocidura f |
| 36 | 40 | 81.6 | 432 | 2 Q7YR10 | Q7Yr10 diceros bic |
| 37 | 40 | 81.6 | 436 | 2 Q7YQM8 | Q7Yqm8 nyctimene a |
| 38 | 40 | 81.6 | 438 | 2 Q7YQM7 | Q7Yqm7 pteropus hy |
| 39 | 40 | 81.6 | 438 | 2 Q7YQR4 | Q7Yqr4 balaena mys |
| 40 | 40 | 81.6 | 438 | 2 Q7YR04 | Q7Yr04 rousettus a |
| 41 | 40 | 81.6 | 441 | 2 Q7YQR3 | Q7Yqr3 phocoenoid |
| 42 | 40 | 81.6 | 443 | 2 Q7YQNS | Q7Yqn5 megaderma l |
| 43 | 40 | 81.6 | 443 | 2 Q7YQP6 | Q7Yqp6 lepus ameri |
| 44 | 40 | 81.6 | 445 | 2 Q7YQNS | Q7Yqn6 bradyus tr |
| 45 | 40 | 81.6 | 445 | 2 Q7YQO7 | Q7Yqo7 tapirus bai |

ALIGNMENTS

RESULT 1
Q7LZ77
ID Q7LZ77 PRELIMINARY; PRT; 275 AA.
AC Q7LZ77;
DT 01-MAR-2004 (TrEMBLrel. 26, Created)
DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DE Apolipoprotein B-100 (Fragment).
OS Gallus Gallus (Chicken).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Archosauria; Aves; Neognathae; Galliformes; Phasianidae;
OC Gallus.
OX NCBI_TaxID=9031;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90324804; PubMed=2373961;
RA Law A., Scott J.;
RT "A cross-species comparison of the apolipoprotein B domain that binds
to the LDL receptor.";
RL J. Lipid Res. 31:1109-1120(1990).
DR PIR; E60950; E60950.
FT NON_TER 1 275
SQ SEQUENCE 275 AA; 30578 MW; B7D8DA054E04B255 CRC64;
Query Match 89.8%; Score 44; DB 2; Length 275;
Best_Local Similarity 90.0%; Pred.No.1.8; Mismatches 1; Indels 0; Gaps 0;
Matches 9; Conservative 0;

Qy 1 TELTRKRGK 10
| | | | | | | | | |
Db 221 TSLTRKRGK 230

RESULT 2
Q7YQN2
ID Q7YQN2 PRELIMINARY; PRT; 387 AA.
AC Q7YQN2;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Phalanger orientalis (gray cuscus).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Metatheria; Diprotodontia; Phalangeridae; Phalanger.
OX NCBI_TaxID=42473;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).

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DR EMBL; AF548431; AAP97387.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 387 387
SQ SEQUENCE 387 AA; 43230 MW; 8300A9D7C54B42B0 CRC64;

Query Match
Best Local Similarity 89.8%; Score 44; DB 2; Length 387;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLK 10
Db 260 TSLTRKRLK 269

RESULT 3
QYQMN PRELIMINARY; PRT; 400 AA.
AC Q7YQMN9;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Ornithorhynchus anatinus (Duckbill platypus).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Monotremata; Ornithorhynchidae; Ornithorhynchus.
OX NCBI_TaxID=9258;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548434; AAP97390.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 400 400
SQ SEQUENCE 400 AA; 44611 MW; DC79873CA6D01CFA CRC64;

Query Match
Best Local Similarity 89.8%; Score 44; DB 2; Length 400;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLK 10
Db 260 TSLTRKRLK 269

RESULT 4
QYQNO PRELIMINARY; PRT; 405 AA.
AC Q7YQNO;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Tachyglossus aculeatus (Australian echidna).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Monotremata; Tachyglossidae; Tachyglossus.
OX NCBI_TaxID=9261;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548433; AAP97389.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 1
```

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FT NON_TER 405 405
SQ SEQUENCE 405 AA; 44975 MW; 551A98557B9B081D CRC64;

Query Match
Best Local Similarity 89.8%; Score 44; DB 2; Length 405;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLK 10
Db 260 TSLTRKRLK 269

RESULT 5
QYQRS PRELIMINARY; PRT; 414 AA.
AC Q7YQRS;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Aotus vociferans (Spix's owl monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Aotus.
OX NCBI_TaxID=57176;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548396; AAP97352.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 414 414
SQ SEQUENCE 414 AA; 45955 MW; EEFA8492157E1BDE CRC64;

Query Match
Best Local Similarity 89.8%; Score 44; DB 2; Length 414;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRLK 10
Db 258 TSLTRKRLK 267

RESULT 6
Q28473 PRELIMINARY; PRT; 596 AA.
AC Q28473;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DE Apolipoprotein B (Fragment).
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopitheidae;
OC Cercopitheinae; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Liver;
RX MEDLINE=92075708; PubMed=1742325;
RA Pape M.E., Castle C.K., Murray R.W., Funk G.M., Hunt C.E.,
RA Marotti K.R., Melchior G.W.;
RT "Apo B metabolism in the cynomolgus monkey: evidence for post-
RT transcriptional regulation.";
RL Biochim. Biophys. Acta 1086:326-334 (1991).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Murray R.;
```

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RL Submitted (FEB-1992) to the EMBL/GenBank/DBSJ databases.
DR EMBL; X15737; CAA33755.1; -.
DR PIR; S32802; S32802.
KW Lipoprotein.
FT NON_TER 1 596
FT NON_TER 596 596
SQ SEQUENCE 596 AA; 66757 MW; B13BBA74E25C3120 CRC64;
Query Match 89.8%; Score 44; DB 2; Length 596;
Best Local Similarity 90.0%; Pred. No. 4;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 226 TELTRKRGK 235

RESULT 7
Q13788 PRELIMINARY; PRT; 3262 AA.
AC Q13788;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE APOB protein (fragment).
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87191999; PubMed=2883086;
RA Carlsson P., Darnfors C., Olofsson S.O., Bjursell G.;
RT "Analysis of the human apolipoprotein B gene: complete structure of
RT the B-74 region.";
RL Gene 49:29-51(1986).
DR EMBL; M15421; AAA51758.1; -.
DR PIR; A27850; LPHUB.
DR GO; GO:0005576; C:extracellular; NAS.
DR GO; GO:0005319; F:lipid transporter activity; NAS.
DR GO; GO:0006869; F:lipid transport; NAS.
FT NON_TER 1 1
SQ SEQUENCE 3262 AA; 370140 MW; 56603BC0618DD40D CRC64;
Query Match 89.8%; Score 44; DB 2; Length 3262;
Best Local Similarity 90.0%; Pred. No. 24;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
Db 2084 TELTRKRGK 2093

RESULT 8
APB HUMAN
ID APB HUMAN STANDARD; PRT; 4563 AA.
AC P04T14; O00502; Q13787;
DT 01-NOV-1986 (Rel. 03, Created)
DT 01-NOV-1986 (Rel. 03, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Apolipoprotein B-100 precursor (Apo B-100) [Contains: Apolipoprotein
DE B-48 (Apo B-48)].
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87016385; PubMed=3763409;
RA Knott T.C., Wallis S.C., Powell L.M., Pease R.J., Lusis A.J.,
RA Blackhart B., McCarthy B.J., Mahley R.W., Levy-Wilson B., Scott J.;
"Complete cDNA and derived protein sequence of human apolipoprotein B-
100.";
Nucleic Acids Res. 14:7501-7503(1986).
[2]
RP SEQUENCE FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=88003974; PubMed=3652907;
RA Ludwig E.H., Blackhart B.D., Pierotti V.R., Caiati L., Fortier C.,
RA Knott T., Scott J., Mahley R.W., Levy-Wilson B., McCarthy B.J.;
RT "DNA sequence of the human apolipoprotein B gene.";
RL DNA 6:363-372(1987).
[3]
RP SEQUENCE FROM N.A., AND VARIANTS ILE-98 AND GLU-4181.
RX MEDLINE=87008488; PubMed=3759943;
RA Chen S.-H., Yang C.-Y., Chen P.-F., Setzer D., Tanimura M., Li W.-H.,
RA Gotto A.M. Jr., Chan L.;
RT "The complete cDNA and amino acid sequence of human apolipoprotein B-
100.";
RL J. Biol. Chem. 261:12918-12921(1986).
[4]
RP SEQUENCE FROM N.A.
RX MEDLINE=87041416; PubMed=3464946;
RA Law S.W., Grant S.M., Higuchi K., Hospattankar A.V., Lackner K.J.,
RA Lee N., Brewer H.B. Jr.;
RT "Human liver apolipoprotein B-100 cDNA: complete nucleic acid and
RT derived amino acid sequence.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:8142-8146(1986).
[5]
RP SEQUENCE FROM N.A.
RX MEDLINE=87161758; PubMed=3030729;
RA Cladaras C., Hadzopoulou-Cladaras M., Nolte R.T., Atkinson D.,
RA Zannis V.I.;
RT "The complete sequence and structural analysis of human apolipoprotein
RT B-100: relationship between apoB-100 and apoB-48 forms.";
RL EMBO J. 5:3495-3507(1986).
[6]
RP SEQUENCE OF 709-906 FROM N.A.
RX MEDLINE=85270450; PubMed=3860836;
RA Deeb S.S., Motulsky A.G., Albers J.J.;
RT "A partial cDNA clone for human apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 82:4983-4986(1985).
[7]
RP SEQUENCE OF 3056-3159 FROM N.A.
RX MEDLINE=86041888; PubMed=3903660;
RA Mehrabian M., Schumaker V.N., Fareed G.C., West R., Johnson D.F.,
RA Kirchessner T.G., Lin H.-C., Wang X., Ma Y., Mendiaz E., Lusis A.J.;
RT "Human apolipoprotein B: identification of cDNA clones and
RT characterization of mRNA.";
Nucleic Acids Res. 13:6937-6953(1985).
[8]
RP SEQUENCE OF 1937-2018 AND 3811-4334 FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=86093680; PubMed=3841204;
RA Carlsson P., Olofsson S.O., Bondjers G., Darnfors C., Wiklund O.,
RA Bjursell G.;
RT "Molecular cloning of human apolipoprotein B cDNA.";
Nucleic Acids Res. 13:8813-8826(1985).
[9]
RP SEQUENCE OF 3109-4563 FROM N.A.
RX MEDLINE=85300528; PubMed=2994225;
RA Knott T.J., Rall S.C. Jr., Imerarity T.L., Jacobson S.F., Urdea M.S.,
RA Levy-Wilson B., Powell L.M., Pease R.J., Eddy R., Nakai H., Byers M.,
RA Priestley L.M., Robertson E., Rall L.B., Betsholtz C., Shows T.B.,
RA Mahley R.W., Scott J.;
RT "Human apolipoprotein B: structure of carboxyl-terminal domains, sites
RT of gene expression, and chromosomal localization.";
Science 230:37-43(1985).
[10]
RP SEQUENCE OF 1-291 FROM N.A.
RX MEDLINE=86149325; PubMed=3513177;
RA Protter A.A., Hardman D.A., Schilling J.W., Miller J., Appleby V.,
RA Chen G.C., Kirshner S.W., McEnroe G., Kane J.P.;
RT "Isolation of a cDNA clone encoding the amino-terminal region of human
RT apolipoprotein B.";
Proc. Natl. Acad. Sci. U.S.A. 83:1467-1471(1986).

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RX SEQUENCE OF 1-1670 FROM N.A., AND VARIANT ILE-98.
 RX MEDLINE=86287319; PubMed=3461454;
 RA Proter A.A., Hardman D.A., Sato K.Y., Schilling J.W., Yamanaka M.,
 RA Hott Y.J., Hjerrild K.A., Chen G.C., Kane J.P.;
 RT "Analysis of cDNA clones encoding the entire B-26 region of human
 RT apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:5678-5682(1986).
 [12]
 RP PARTIAL SEQUENCE, AND IDENTIFICATION OF APO-B48.
 RX MEDLINE=88018019; PubMed=3659919;
 RA Chen S.-H., Habib G., Yang C.-H., Gu Z.-W., Lee B.R., Weng S.-H.,
 RA Silbermann S.R., Cai S.-J., Deslypere J.P., Rosseneu M.,
 RA Gotto A.M. Jr., Li W.-H., Chan L.;
 RT "Apolipoprotein B-48 is the product of a messenger RNA with an organ-
 RT specific in-frame stop codon.";
 RL Science 238:363-366(1987).
 [13]
 RP DOMAINS.
 RX MEDLINE=87039351; PubMed=3773997;
 RA Knott T.C., Pease R.J., Powell L.M., Wallis S.C., Rall S.C. Jr.,
 RA Innerarity T.L., Blackhart B., Taylor W.R., Marcel Y., Milne R.,
 RA Johnson D., Fuller M., Lusis A.J., McCarthy B.J., Mahley R.W.,
 RA Levy-Wilson B., Scott J.;
 RT "Complete protein sequence and identification of structural domains of
 RT human apolipoprotein B.";
 RL Nature 323:734-738(1986).
 [14]
 RP DOMAINS.
 RX MEDLINE=86242245; PubMed=3087360;
 RA Daehtli N., Lee D.M., Mok T.;
 RA "Apolipoprotein B is a calcium binding protein.";
 RL Biochem. Biophys. Res. Commun. 137:493-499(1986).
 [16]
 RP PALMITOYLATION OF CYS-1112.
 RX MEDLINE=20143590; PubMed=10679026;
 RA Zhao Y., McCabe J.B., Vance J., Berthiaume L.G.;
 RT "Palmitoylation of apolipoprotein B is required for proper
 RT intracellular sorting and transport of cholesterol esters and
 RT triglycerides.";
 RL Mol. Biol. Cell 11:721-734(2000).
 [17]
 RP VARIANT SER-4338.
 RX MEDLINE=91071750; PubMed=1979313;
 RA Navaajas M., Laurent A.-M., Moreel J.-F., Ragab A., Cambou J.-P.,
 RA Cuny G., Cambien F., Roizes G.;
 RT "Detection by denaturing gradient gel electrophoresis of a new
 RT polymorphism in the apolipoprotein B gene.";
 RL Hum. Genet. 86:91-93(1990).
 [18]
 RP VARIANT FDB GLN-3527.
 RX MEDLINE=89098975; PubMed=2563166;
 RA Soria L.F., Ludwig E.H., Clarke H.R.G., Vega G.L., Grundy S.M.,
 RA McCarthy B.J.;
 RT "Association between a specific apolipoprotein B mutation and familial
 RT defective apolipoprotein B-100.";
 RL Proc. Natl. Acad. Sci. U.S.A. 86:587-591(1989).
 [19]
 RP VARIANT LEU-2739.
 RX MEDLINE=91016974; PubMed=2216805;
 RA Huang L.-S., Gavish D., Breslow J.L.;
 RT "Sequence polymorphism in the human apoB gene at position 8344.";
 RL Nucleic Acids Res. 18:5922-5922(1990).
 [20]
 RP VARIANT FDB CYS-3558.

RX MEDLINE=95190020; PubMed=7893971;
 RA Pullinger C.R., Hennessy L.K., Chatterton J.E., Liu W., Love J.A.,
 RA Mendel C.M., Frost P.H., Malloy M.J., Schumaker V.N., Kane J.P.;
 RT "Familial ligand-defective apolipoprotein B. Identification of a new
 RT mutation that decreases LDL receptor binding affinity.";
 RL J. Clin. Invest. 95:1225-1234(1995).
 [21]
 RP VARIANTS LEU-1437; SER-1914; LYS-2566; THR-3121; ALA-3945; MET-4128
 RP AND THR-4481.
 RX MEDLINE=97044521; PubMed=8889592;
 RA Poirier O., Ricard S., Behague I., Souriau C., Evans A.E.,
 RA Arveiler D., Marques-Vidal P., Luc G., Roizes G., Cambien F.;
 RT "Detection of new variants in the apolipoprotein B (Apo B) gene by
 RT PCR-SSCP.";
 RL Hum. Mutat. 8:282-285(1996).
 [22]
 RP VARIANTS FDB GLN-3527 AND CYS-3558.
 RX MEDLINE=97403938; PubMed=9259199;
 RA Rabes J.P., Varret M., Saint-Jore B., Erlich D., Jondeau G.,
 RA Krempf M., Giraudeau P., Junien C., Boileau C.;
 RT "Familial ligand-defective apolipoprotein B-100: simultaneous
 RT detection of the ARG3500-->GLN and ARG3531-->CYS mutations in a French
 RT population.";
 RL Hum. Mutat. 10:160-163(1997).
 [23]
 RP VARIANTS SER-1914; ARG-1923; LEU-2739; ASP-3319; THR-3427; GLN-3432
 RP AND ILE-3921.
 RX MEDLINE=98141125; PubMed=9490296;
 RA Leren T.P., Bakken K.S., Hoel V., Hjermann I., Berg K.;
 RT "Screening for mutations of the apolipoprotein B gene causing
 RT hypocholesterolemia.";
 RL Hum. Genet. 102:44-49(1998).
 CC -!- FUNCTION: Apolipoprotein B is a major protein constituent of
 CC chylomicrons (apo B-48), LDL (apo B-100) and VLDL (apo B-100). Apo
 CC B-100 functions as a recognition signal for the cellular binding
 CC and internalization of LDL particles by the apoB/E receptor.
 CC -!- SUBCELLULAR LOCATION: Secreted.

Query Match 89.8%; Score 44; DB 1; Length 4563;
 Best Local Similarity 90.0%; Pred. No. 35;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
 Db 3385 TELTRKRGK 3394

RESULT 9
 Q7Z600
 ID Q7Z600 PRELIMINARY; PRT; 4563 AA.
 AC Q7Z600;
 DT 01-OCT-2003 (TREMELrel. 25, Created)
 DT 01-OCT-2003 (TREMELrel. 25, Last sequence update)
 DT 01-MAR-2004 (TREMELrel. 26, Last annotation update)
 DE Apolipoprotein B (Including Ag(X) antigen).
 GN Name=APOB;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.
 RN [1]_NCBI_TaxID=9606;
 RP SEQUENCE FROM N.A.
 RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
 RA Ahearn M.O., Kuldanek S.A., Rajkumar N., Toth E.J., Yi Q.,
 RA Nickerson D.A.;
 RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY324608; AAP72970.1; -.
 DR GO; GO:0005319; F:lipid transporter activity; IEA.
 DR GO; GO:0006869; P:lipid transport; IEA.
 DR InterPro; IPR009454; DUF1081.
 DR InterPro; IPR001747; Lipid transprt_N.
 DR Pfam; PF06448; DUF1081; 1.
 DR Pfam; PF01347; Vitellogenin_N; 1.


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DR SMART; SM00638; LPD_N; 1.
KW Lipoprotein.
SQ SEQUENCE 4563 AA; 515553 MW; 030B34167CDBC63C CRC64;

Query Match
Best Local Similarity 89.8%; Score 44; DB 2; Length 4563;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
DB 3385 TELTRKRGK 3394

RESULT 10
Q6PIM8
ID Q6PIM8 PRELIMINARY; PRT; 164 AA.
AC Q6PIM8;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DE ACSL3 protein (Fragment).
GN Name=ACSL3;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L., Feingold E.A., Grouse L.H., Derge J.G.,
RA Klausner R.D., Collins F.S., Wagner L., Shenmen C.M., Schuler G.D.,
RA Altschul S.F., Zeeberg B., Buetow K.H., Schaefer C.F., Bhat N.K.,
RA Hopkins R.F., Jordan H., Moore T., Max S.I., Wang J., Hsieh F.,
RA Diatchenko L., Marusina K., Farmer A.A., Rubin G.M., Hong L.,
RA Stapleton M., Soares M.B., Bonaldo M.F., Casavant T.L., Scheetz T.E.,
RA Brownstein M.J., Usdin T.B., Toshiyuki S., Carninci P., Prange C.,
RA Raha S.S., Loquellano N.A., Peters G.J., Abramson R.D., Mullahy S.J.,
RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Villalon D.K., Muzny D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodriguez S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzywinski M.I., Skalska U., Smailus D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Lung;
RA Strausberg R.;
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; BC032144; AAH32144.1; -.
FT NON TER 1
SQ SEQUENCE 164 AA; 18619 MW; 0BB00FE1A649E9AA CRC64;

Query Match 83.7%; Score 41; DB 2; Length 164;
Best Local Similarity 80.0%; Pred. NO. 4.2;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
DB 77 TELARKKGLK 86

RESULT 12
LCF3 HUMAN
ID _LCF3_HUMAN STANDARD; PRT; 720 AA.
AC 095573;
DT 16-OCT-2001 (Rel. 40, Created)
DT 16-OCT-2001 (Rel. 40, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Long-chain-fatty-acid--CoA ligase 3 (EC 6.2.1.3) (Long-chain acyl-CoA
DE synthetase 3) (LACS 3).
GN Name=ACSL3; Synonyms=FACL3, ACS3, LACS3;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=97321062; PubMed=9177793;
RA Minekura H., Fujino T., Kang M.-J., Fujita T., Endo Y., Yamamoto T.T.;
RT "Human acyl-coenzyme A synthetase 3 cDNA and localization of its gene
RT (ACS3) to chromosome band 2q34-q35.";
RL Genomics 42:180-181(1997).
RN [2]

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SEQUENCE FROM N.A.
RX MEDLINE=21564184; PubMed=11707336; DOI=10.1016/S0378-1119(01)00714-4;
RA Minekura H., Kang M.-J., Inagaki Y., Suzuki H., Sato H., Fujino T.,
RA Yamamoto T.T.;
RT "Genomic organization and transcription units of the human acyl-CoA
RL Gene 278:185-192(2001).
CC -!- FUNCTION: Activation of long-chain fatty acids for both synthesis
CC of cellular lipids, and degradation via beta-oxidation.
CC Preferentially uses myristate, laurate, arachidonate and
CC eicosapentaenoate as substrates (By similarity).
CC -!- CATALYTIC ACTIVITY: ATP + a long-chain carboxylic acid + CoA = AMP
CC + diphosphate + an acyl-CoA.
CC -!- COFACTOR: Magnesium (By similarity).
CC -!- SUBCELLULAR LOCATION: Mitochondria; outer mitochondrial membrane and
CC peroxisomal membrane.
CC -!- SIMILARITY: Belongs to the ATP-dependent AMP-binding enzyme
CC family.
CC
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC entities requires a license agreement (See http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC
CC -----
CC EMBL; D89053; BAA37142.1; -.
CC EMBL; AB061712; BAB72074.1; -.
CC GenBank; HGNC:3570; ACSL3.
CC MIM; 602371; -.
CC GO; GO:0004321; F:fatty-acyl-CoA synthase activity; TAS.
CC InterPro; IPR000873; AMP-bind.
CC Pfam; PF00501; AMP-binding; 1.
CC PRINTS; PR00154; AMPBINDING.
CC PROSITE; PS00455; AMP BINDING; 1.
CC Fatty acid metabolism; Ligase; Magnesium; Multigene family.
CC SEQUENCE 720 AA; 80345 MW; 845959A765BC6BF6 CRC64;

Query Match 83.7%; Score 41; DB 1; Length 720;
Best Local Similarity 80.0%; Pred. No. 20;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
DB |||||:||||
633 TELARKKGLK 642

RESULT 13
Q8IU09 PRELIMINARY; PRT; 720 AA.
AC O8IU09;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DE 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Acyl-CoA synthetase long-chain family member 3.
GN Name=ACSL3;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Minekura H., Kang M.-J., Inagaki Y., Suzuki H., Sato H., Fujino T.,
RA Yamamoto T.T.;
RT "Genomic organization and transcription units of the human acyl-CoA
RL Gene 278:185-192(2001).
CC EMBL; AB061436; BAB72139.1; -.
CC EMBL; AB061423; BAB72139.1; JOINED.
CC EMBL; AB061424; BAB72139.1; JOINED.
CC EMBL; AB061425; BAB72139.1; JOINED.
CC EMBL; AB061426; BAB72139.1; JOINED.
CC EMBL; AB061427; BAB72139.1; JOINED.
CC EMBL; AB061428; BAB72139.1; JOINED.
CC EMBL; AB061429; BAB72139.1; JOINED.
CC EMBL; AB061430; BAB72139.1; JOINED.
CC EMBL; AB061431; BAB72139.1; JOINED.
CC EMBL; AB061432; BAB72139.1; JOINED.
CC EMBL; AB061433; BAB72139.1; JOINED.
CC EMBL; AB061434; BAB72139.1; JOINED.
CC EMBL; AB061435; BAB72139.1; JOINED.
CC SEQUENCE 720 AA; 80345 MW; 845959A765BC6BF6 CRC64;

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RA Bosak S.A., McEwan P.J., McKernan K.J., Malek J.A., Gunaratne P.H.,
RA Richards S., Worley K.C., Hale S., Garcia A.M., Gay L.J., Hulyk S.W.,
RA Vallalon D.K., Munzy D.M., Sodergren E.J., Lu X., Gibbs R.A.,
RA Fahey J., Helton E., Kettman M., Madan A., Rodrigues S., Sanchez A.,
RA Whiting M., Madan A., Young A.C., Shevchenko Y., Bouffard G.G.,
RA Blakesley R.W., Touchman J.W., Green E.D., Dickson M.C.,
RA Rodriguez A.C., Grimwood J., Schmutz J., Myers R.M., Butterfield Y.S.,
RA Krzyzinski M.I., Skalska U., Smailus D.E., Schnerch A., Schein J.E.,
RA Jones S.J., Marra M.A.;
RA "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences."
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Pancreas;
RA Strausberg R.;
RL Submitted (DEC-2002) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: Belongs to the ATP-dependent AMP-binding enzyme
CC family.
CC EMBL; BC041692; AAH41692.1; -.
CC GO; GO:0003824; F:catalytic activity; IEA.
CC GO; GO:0008152; P:metabolism; IEA.
CC InterPro; IPR000873; AMP-bind.
CC Pfam; PF00501; AMP-binding; 1.
CC PRINTS; PR00154; AMPBINDING.
CC PROSITE; PS00455; AMP BINDING; 1.
CC SEQUENCE 720 AA; 80419 MW; AAC4B0B4543EC8DD CRC64;

Query Match 83.7%; Score 41; DB 2; Length 720;
Best Local Similarity 80.0%; Pred. No. 20;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TELTRKRGK 10
DB |||||:||||
633 TELARKKGLK 642

RESULT 14
BAB72139 PRELIMINARY; PRT; 720 AA.
AC BAB72139;
DT 02-MAR-2004 (TrEMBLrel. 27, Created)
DT 02-MAR-2004 (TrEMBLrel. 27, Last sequence update)
DE 02-MAR-2004 (TrEMBLrel. 27, Last annotation update)
DE Acyl-CoA synthetase 3.
GN FACL3.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RA Minekura H., Kang M.-J., Inagaki Y., Suzuki H., Sato H., Fujino T.,
RA Yamamoto T.T.;
RT "Genomic organization and transcription units of the human acyl-CoA
RL Gene 278:185-192(2001).
CC EMBL; AB061436; BAB72139.1; -.
CC EMBL; AB061423; BAB72139.1; JOINED.
CC EMBL; AB061424; BAB72139.1; JOINED.
CC EMBL; AB061425; BAB72139.1; JOINED.
CC EMBL; AB061426; BAB72139.1; JOINED.
CC EMBL; AB061427; BAB72139.1; JOINED.
CC EMBL; AB061428; BAB72139.1; JOINED.
CC EMBL; AB061429; BAB72139.1; JOINED.
CC EMBL; AB061430; BAB72139.1; JOINED.
CC EMBL; AB061431; BAB72139.1; JOINED.
CC EMBL; AB061432; BAB72139.1; JOINED.
CC EMBL; AB061433; BAB72139.1; JOINED.
CC EMBL; AB061434; BAB72139.1; JOINED.
CC EMBL; AB061435; BAB72139.1; JOINED.
CC SEQUENCE 720 AA; 80345 MW; 845959A765BC6BF6 CRC64;

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Query Match 83.7%; Score 41; DB 2; Length 720;
Best Local Similarity 80.0%; Pred.No. 20;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLK 10
Db 633 TELARKKGLK 642

RESULT 15

ID Q7YQO0 PRELIMINARY; PRT; 263 AA.
AC Q7YQO0;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Procavia capensis (Cape hyrax) (Rock dassie).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Hyracoidea; Procaviidae; Procavia.
OX NCBI_TaxID=9813;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548411; AAP97367.1; -.
KW Lipoprotein.
FT NON TER 1
FT NON TER 263
SQ SEQUENCE 263 AA; 29532 MW; 536CF6149C1D062A CRC64;

Query Match 81.6%; Score 40; DB 2; Length 263;
Best Local Similarity 80.0%; Pred.No. 11;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TELTRKRLK 10
Db 206 SSLTRKRLK 215

Search completed: January 13, 2005, 01:51:02
Job time : 78.0328 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 77.2131 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-12
Perfect score: 50
Sequence: 1 TDLTRKRLK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_23Sep04:*

- 1: geneseqp1980s:*
- 2: geneseqp1990s:*
- 3: geneseqp2000s:*
- 4: geneseqp2001s:*
- 5: geneseqp2002s:*
- 6: geneseqp2003as:*
- 7: geneseqp2003bs:*
- 8: geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB | ID | Description |
|------------|-------|-------------|--------|----|----------|--------------------|
| 1 | 50 | 100.0 | 10 | 2 | AAY30693 | Aay30693 Apo-B100 |
| 2 | 46 | 92.0 | 10 | 2 | AAY30692 | Aay30692 Apo-B100 |
| 3 | 42 | 84.0 | 11 | 2 | Aaw57205 | Aaw57205 Apo B bin |
| 4 | 42 | 84.0 | 13 | 2 | Aaw57207 | Aaw57207 Apo B 100 |
| 5 | 42 | 84.0 | 15 | 2 | Aaw41261 | Aaw41261 Apolipop |
| 6 | 42 | 84.0 | 15 | 2 | Aaw96892 | Abj37575 Heparin b |
| 7 | 42 | 84.0 | 20 | 6 | ABJ37575 | Aaw57208 Apo B 100 |
| 8 | 42 | 84.0 | 22 | 2 | Aaw57209 | Aaw57209 Apo B 100 |
| 9 | 42 | 84.0 | 34 | 5 | Aae14541 | Aae14541 Human apo |
| 10 | 42 | 84.0 | 36 | 2 | Aaw96876 | Aaw96876 Nucleic a |
| 11 | 42 | 84.0 | 37 | 2 | Aaw64587 | Aaw64587 Human apo |
| 12 | 42 | 84.0 | 51 | 2 | Aaw96845 | Aaw96845 Nucleic a |
| 13 | 42 | 84.0 | 343 | 4 | ABB37687 | Abb37687 Peptide # |
| 14 | 42 | 84.0 | 343 | 4 | ABG52504 | Abg52504 Human liv |
| 15 | 42 | 84.0 | 377 | 2 | AAR72704 | Aar72704 Human apo |
| 16 | 42 | 84.0 | 377 | 2 | AAR34031 | Aar34031 Sequence |
| 17 | 42 | 84.0 | 2463 | 8 | ADJ57400 | Adj57400 Human apo |
| 18 | 42 | 84.0 | 3923 | 2 | AAY31237 | Aay31237 Human Apo |
| 19 | 42 | 84.0 | 4536 | 2 | Aaw41262 | Aaw41262 Apolipop |
| 20 | 42 | 84.0 | 4536 | 2 | Aaw96826 | Aaw96826 Amino aci |
| 21 | 42 | 84.0 | 4560 | 5 | Aau98981 | Aau98981 Human apo |
| 22 | 42 | 84.0 | 4561 | 7 | ADD48677 | Add48677 Human pro |
| 23 | 42 | 84.0 | 4563 | 5 | AAO15893 | Aao15893 Human apo |
| 24 | 42 | 84.0 | 4563 | 6 | ABR40253 | AbR40253 Human ali |
| 25 | 42 | 84.0 | 4563 | 6 | ABR40253 | AbR40253 Human ali |

| | | | | | | |
|----|----|------|------|---|-----------|--------------------|
| 26 | 42 | 84.0 | 4563 | 6 | ABU79140 | Abu79140 Apolipop |
| 27 | 42 | 84.0 | 4563 | 7 | ADF43408 | Adf43408 Apolipop |
| 28 | 42 | 84.0 | 4563 | 8 | ADH18871 | Adh18871 Human apo |
| 29 | 42 | 84.0 | 4563 | 8 | ADH18870 | Adh18870 Human apo |
| 30 | 42 | 84.0 | 4563 | 8 | ADO33445 | Ado33445 Human apo |
| 31 | 42 | 84.0 | 4563 | 8 | ADO33447 | Ado33447 Human apo |
| 32 | 42 | 84.0 | 4590 | 4 | AAU33184 | Aau33184 Novel hum |
| 33 | 40 | 80.0 | 782 | 7 | ADA06278 | Ada06278 Human cel |
| 34 | 39 | 78.0 | 11 | 2 | AAW57206 | Aaw57206 Apo B 100 |
| 35 | 39 | 78.0 | 11 | 2 | AAW87717 | Aaw87717 Analogue |
| 36 | 39 | 78.0 | 11 | 5 | AAE21732 | Aae21732 BSMR effe |
| 37 | 39 | 78.0 | 11 | 6 | ABU07938 | Abu07938 Apoprotei |
| 38 | 39 | 78.0 | 11 | 7 | ADF56451 | Adf56451 Human apo |
| 39 | 39 | 78.0 | 12 | 2 | AAW41260 | Aaw41260 Apolipop |
| 40 | 39 | 78.0 | 15 | 2 | AAW22911 | Aaw22911 Low densi |
| 41 | 39 | 78.0 | 23 | 6 | ABR57177 | AbR57177 Human PDG |
| 42 | 38 | 76.0 | 10 | 2 | AAAY30682 | Aay30682 Apo-B100 |
| 43 | 38 | 76.0 | 10 | 2 | AAAY30687 | Aay30687 Apo-B100 |
| 44 | 38 | 76.0 | 63 | 4 | ABG09607 | Abg09607 Novel hum |
| 45 | 38 | 76.0 | 465 | 4 | AAB92994 | Aab92994 Human pro |

ALIGNMENTS

RESULT 1
AAY30693
ID AAY30693 standard; peptide; 10 AA.
XX
AC AAY30693;
XX
DT 17-NOV-1999 (first entry)
XX
DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS Synthetic.
OS Homo sapiens.
XX
FN WO9946598-A1.
XX
PD 16-SEP-1999.
XX
PF 05-MAR-1999; 99WO-US004805.
XX
PR 10-MAR-1998; 98US-0077618P.
XX (REGC) UNIV CALIFORNIA.
XX
PI Innerarity TL, Boren JOS;
XX
DR WPI; 1999-551509/46.
XX
PT Identifying compounds which affect binding of low density lipoprotein
PT with proteoglycan, used for, e.g. obtaining compounds for reducing
PT atherosclerosis.
XX
PS Claim 17; Page 57; 70pp; English.
XX
CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC receptor mutations. They were created to identify compounds which
CC modulate atherosclerosis. The peptides are derived from amino acids 3358
CC to 3367 of apoB100. The method comprises detecting compounds which affect
CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC can be used for identifying compounds which disrupt LDL-PG binding
CC without inhibiting LDL receptor binding. Such compounds can be used to
CC reduce or prevent the formation of atherosclerotic lesions and prevent
CC atherosclerosis. The transgenic non-human animals and mammals which
CC express human apo-B100 can be used as an in vivo model system for the
CC study of atherosclerosis, and in vivo assay methods for identifying
CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

XX SQ Sequence 10 AA;

Query Match 100.0%; Score 50; DB 2; Length 10;
 Best Local Similarity 100.0%; Pred. No. 0.0062;
 Matches 10; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
 |:|||||
 Db 1 TDLTRKRGGLK 10

RESULT 2

AAV30692

ID AAY30692 standard; peptide; 10 AA.
 XX
 AC AAY30692;
 XX 17-NOV-1999 (first entry)
 XX Apo-B100 derived peptide showing a proteoglycan receptor mutation.

XX Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

XX Synthetic.

OS Homo sapiens.

XX WO9946598-A1.

FN 16-SEP-1999.

XX 05-MAR-1999; 99WO-US004805.

XX 10-MAR-1998; 98US-0077618P.

XX (REGC) UNIV CALIFORNIA.

PA Innerarity TL, Boren JOS;

XX WPI; 1999-551509/46.

XX Identifying compounds which affect binding of low density lipoprotein
 CC with proteoglycan, used for, e.g. obtaining compounds for reducing
 CC atherosclerosis.

PS Claim 17; Page 57; 70pp; English.

XX AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 10 AA;

Query Match 92.0%; Score 46; DB 2; Length 10;
 Best Local Similarity 90.0%; Pred. No. 0.038;
 Matches 9; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
 |:|||||
 Db 1 TELTRKRGGLK 10

RESULT 3

AAW57205

ID AAW57205 standard; peptide; 11 AA.
 XX
 AC AAW57205;
 XX 03-AUG-1998 (first entry)
 XX Apo B binding site peptide 2.
 XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.

XX Synthetic.

XX WO9813385-A2.

XX 02-APR-1998.

XX 25-SEP-1997; 97WO-GB002610.

XX 27-SEP-1996; 96GB-00020153.

XX (UYST) UNIV STRATHCLYDE.

XX Halbert GW, Owens MD, Baillie G;

XX WPI; 1998-230637/20.

XX Non-natural lipid particle comprising peptide binding to apo B protein
 CC receptor - useful as, e.g. vector for delivering drugs to cancer cells
 CC that express this receptor.

PS Claim 12; Page 52; 73pp; English.

XX The present sequence represents a specifically claimed Apo B binding site
 CC peptide which can be used as a component of a non-naturally occurring,
 CC receptor-competent low density lipoprotein (LDL) particle of the present
 CC invention. The LDL particle comprises at least 1 peptide component that
 CC has at least 1 binding site for an apo B protein receptor and at least 1
 CC lipophilic substituent. Also described in the invention are peptides
 CC containing an apo B binding sequence with at least 70% identity with
 CC sequences: KAVYKMKHRH (1) or TRLTRKRGGLK (2), or their dimers. Non-
 CC naturally occurring, receptor-competent LDL particles are useful as: (i)
 CC drug-targeting vectors for delivering anticancer drugs to cancer cells
 CC that express an apo B protein receptor, and (ii) additives for cell
 CC culture media especially as growth supplements. Non-naturally occurring,
 CC receptor-competent LDL particles do not require the complete apo B
 CC sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor

XX SQ Sequence 11 AA;

Query Match 84.0%; Score 42; DB 2; Length 11;
 Best Local Similarity 90.0%; Pred. No. 0.25;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
 |:|||||
 Db 2 TELTRKRGGLK 11

RESULT 4
 AAW57207
 ID AAW57207 standard; peptide; 13 AA.
 XX
 AC AAW57207;
 XX
 DT 03-AUG-1998 (first entry)
 XX
 DE Apo B 100 binding site peptide analogue peptide B.
 XX
 KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 OS Synthetic.
 OS
 XX
 FH Key Location/Qualifiers
 FT Modified-site 1 /note= "attached to retinoic acid"
 FT
 XX WO9813385-A2.
 XX
 XX 02-APR-1998.
 PD
 XX
 PF 25-SEP-1997; 97WO-GB002610.
 XX
 XX 27-SEP-1996; 96GB-00020153.
 PR
 XX (UYST) UNIV STRATHCLYDE.
 PA
 XX
 PI Halbert GW, Owens MD, Baillie G;
 XX
 XX WPI; 1998-230637/20.
 DR
 XX
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 PT
 XX
 PS Claim 13; Fig 7; 73pp; English.
 XX
 CC The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRGK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 13 AA;
 Query Match 84.0%; Score 42; DB 2; Length 13;
 Best Local Similarity 90.0%; Pred. No. 0.3;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TDLTRKRGK 10
 | | | | | | | |
 Db 3 TRLTRKRGK 12
 RESULT 5
 AAW41261
 ID AAW41261 standard; peptide; 15 AA.
 XX
 AC AAW41261;

XX 19-MAY-1998 (first entry)
 DT
 XX Apolipoprotein B-100 fragment.
 DE
 XX
 KW Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
 KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
 KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
 KW prothrombinase complex.
 XX
 OS Synthetic.
 OS
 OS Homo sapiens.
 XX
 PN WO9743311-A1.
 XX
 XX 20-NOV-1997.
 PD
 XX
 XX 09-MAY-1997; 97WO-GB001255.
 PF
 XX 09-MAY-1996; 96GB-00009702.
 PR
 XX (UNLO) ROYAL FREE HOSPITAL SCHOOL MED.
 PA
 XX Bruckdorfer KR, Etelaie C;
 XX
 PI WPI; 1998-008798/01.
 DR
 XX
 XX Peptide fragments of apo:lipo:protein B-100 with anticoagulant activity -
 PT used for treating or preventing coagulation, inhibiting angiogenesis,
 PT cell differentiation and apoptosis.
 PT
 XX Disclosure; Page 22; 60pp; English.
 PS
 XX
 CC This sequence is an example of the peptide of the invention. It has the
 CC formula (I), or their variants with one or more internal deletions,
 CC insertions or substitutions, while retaining anti-coagulant properties of
 CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KKNKRRHS-X2-T-22 (I) X1 = S or
 CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
 CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
 CC aa. Compositions containing the peptide are used for simultaneous,
 CC separate or sequential treatment of cancer, particularly to prevent
 CC metastatic spread. They are also used to inhibit thromboplastin-mediated
 CC processes, specifically to prevent or reduce blood coagulation (e.g.
 CC during or after surgery or in cases of heart attack, stroke etc.) and to
 CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
 CC which is active as such or as part of a 98-aa peptide, inhibits
 CC activation of the prothrombinase complex; and prevents activation of
 CC factor VII on the surface of thromboplastin and of platelets by thrombin.
 CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
 CC smaller than apoB-100, they act more quickly
 XX
 SQ Sequence 15 AA;
 Query Match 84.0%; Score 42; DB 2; Length 15;
 Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TDLTRKRGK 10
 | | | | | | | |
 Db 1 TRLTRKRGK 10
 RESULT 6
 AAW96892
 ID AAW96892 standard; peptide; 15 AA.
 XX
 AC AAW96892;
 XX
 XX 22-APR-1999 (first entry)
 DT
 XX ApoB-100 nuclear localisation signal sequence, residues 3363-3367.
 DE
 XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;

KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.

XX Homo sapiens.

PN WO9856938-A1.

XX 17-DEC-1998.

XX 10-JUN-1998; 98WO-US011927.

XX 13-JUN-1997; 97US-00874807.

PR 14-MAY-1998; 98US-00079030.

XX (BAYU) BAYLOR COLLEGE MEDICINE.

XX Guevara JG, Hoogeveen RC, Moore JP;

XX WPI; 1999-070331/06.

XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.

XX Claim 19; Fig 13D; 293pp; English.

XX AAW96878-97 represent nuclear localisation signal sequence derived from
 CC human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein
 CC component of very-low density lipoproteins (VLDL), intermediate density
 CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The
 CC present sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis

XX SQ Sequence 15 AA;

Query Match 84.0%; Score 42; DB 2; Length 15;

Best Local Similarity 90.0%; Pred. No. 0.34;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRGK 10
 | | | | |
 Db 6 TRLTRKRGK 15

RESULT 7

ABJ37575
 ID ABJ37575 standard; peptide; 20 AA.

XX AC ABJ37575;

XX DT 10-MAY-2003 (first entry)

XX Heparin binding peptide sequence #28.

XX Cytostatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;
 KW cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;
 KW rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.

XX Unidentified.

XX WO2003007689-A2.

XX 30-JAN-2003.

XX 22-JUL-2002; 2002WO-US023419.

XX

PR 20-JUL-2001; 2001US-0306726P.

XX (ETHZ-) ETH ZUERICH.

PA (UYZU-) UNIV ZURICH.

XX Hubbell JA, Schoenmakers R, Maynard HD;

XX WPI; 2003-300420/29.

XX Use of a ligand comprising of at least one sulfated or sulfonated amino
 PT acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic
 PT retinopathy and hypoxia.

XX Disclosure; Fig 2; 79pp; English.

XX The invention relates to a novel ligand for binding a target biomolecule,
 CC which comprises a peptide having at least one sulphated or sulphonated
 CC amino acid and at least one amino acid chosen from neutral and positively
 CC charged amino acids. The novel ligands can be used for the treatment of
 CC e.g. tumours, rheumatoid arthritis, diabetic retinopathy and hypoxia.
 CC This sequence represents a heparin binding peptide relating to the
 CC invention

XX SQ Sequence 20 AA;

Query Match 84.0%; Score 42; DB 6; Length 20;

Best Local Similarity 90.0%; Pred. No. 0.46;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRGK 10
 | | | | |
 Db 7 TRLTRKRGK 16

RESULT 8

AAW57208

ID AAW57208 standard; peptide; 22 AA.

XX AC AAW57208;

XX DT 03-AUG-1998 (first entry)

XX Apo B 100 binding site peptide analogue peptide C.

XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.

XX Synthetic.

XX Key Location/Qualifiers

FT Modified-site 1 /note= "attached to retinoic acid"

FT Modified-site 22 /note= "attached to cholesterol"

FT Modified-site 22 /note= "attached to cholesterol"

FT Modified-site 22 /note= "attached to cholesterol"

XX WO9813385-A2.

XX 02-APR-1998.

XX 25-SEP-1997; 97WO-GB002610.

XX 27-SEP-1996; 96GB-00020153.

XX (UYST) UNIV STRATHCLYDE.

XX Halbert GW, Owens MD, Baillie G;

XX WPI; 1998-230637/20.

XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.

XX Claim 13; Fig 7; 73pp; English.
 XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 22 AA;

Query Match 84.0%; Score 42; DB 2; Length 22;
 Best Local Similarity 90.0%; Pred. No. 0.5;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLK 10
 | | | | |
 Db 7 TRLTRKRLK 16

RESULT 9
 AAWS7209
 ID AAW57209 standard; peptide; 22 AA.
 AC AAW57209;
 XX
 DT 03-AUG-1998 (first entry)
 XX
 DE Apo B 100 binding site peptide analogue peptide D.
 XX
 KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 OS Synthetic.

Key Location/Qualifiers
 FT Modified-site 1 /note= "attached to retinoic acid"
 FT

XX WO9813385-A2.
 XX
 PD 02-APR-1998.
 XX
 PF 25-SEP-1997; 97WO-GB002610.
 XX
 PR 27-SEP-1996; 96GB-00020153.
 XX
 PA (UVST) UNIV STRATHCLYDE.

XX Halbert GW, Owens MD, Baillie G;
 XX
 DR WPI; 1998-230637/20.

XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.

XX Claim 13; Fig 7; 73pp; English.

XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)

CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 22 AA;

Query Match 84.0%; Score 42; DB 2; Length 22;
 Best Local Similarity 90.0%; Pred. No. 0.5;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLK 10
 | | | | |
 Db 7 TRLTRKRLK 16

RESULT 10
 AAET14541
 ID AAE14541 standard; peptide; 34 AA.
 XX
 AC AAE14541;
 XX

DT 17-MAY-2002 (first entry)
 XX
 DE Human apoB-100 derived peptide p62.

XX Oxidised low density lipoprotein; oxLDL; immunoassay; periaortitis;
 KW cardiovascular disease; coronary heart disease; pre-eclampsia;
 KW non-insulin-dependent diabetes; endothelial dysfunction; human; apoB-100;
 KW peptide p62.

XX Homo sapiens.

XX WO200206314-A2.

XX 24-JAN-2002.

XX 18-JUL-2001; 2001WO-GB003212.

XX 18-JUL-2000; 2000GB-00017641.

XX (ARKT-) ARK THERAPEUTICS LTD.

XX Narvanen O, Yla-Herttuala S;

XX WPI; 2002-179777/23.

XX New peptide useful in enzyme immunoassays for detecting oxidized low
 PT density lipoprotein which is a marker of coronary heart disease and other
 PT cardiovascular diseases, has affinity for oxidized low density
 PT lipoprotein.

XX Claim 6; Page 5; 21pp; English.

XX The invention relates to peptides having affinity for oxidised low
 CC density lipoprotein (oxLDL), in cyclised or multimeric form. The peptide
 CC is useful in an immunoassay to determine the presence, and optionally,
 CC the amount of antibodies in a sample, having affinity for oxLDL.
 CC Preferably immobilised peptide is useful for measuring the amount of
 CC autoantibodies for oxLDL in a sample, especially a serum or plasma sample
 CC from a patient for evaluating the risk of coronary heart diseases, other
 CC cardiovascular diseases, and several other disorders such as
 CC periaortitis, pre-eclampsia, non-insulin-dependent diabetes and
 CC endothelial dysfunction. The peptide of the invention is stable, can be
 CC synthesised easily without the need to isolate proteins from a patient's

CC blood, and has a long half-life. The present sequence is human apoB-100
 CC derived peptide p62 used in the invention
 XX
 SQ Sequence 34 AA;

Query Match 84.0%; Score 42; DB 5; Length 34;
 Best Local Similarity 90.0%; Pred. No. 0.78;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRLK 10
 | | | | |
 DB 25 TRLTRKRLK 34

RESULT 11
 AAW96876
 ID AAW96876 standard; peptide; 36 AA.

XX AAW96876;
 AC
 DT 22-APR-1999 (first entry)
 DE Nucleic acid binding domain from apoB-100, residues 3348-3390.
 XX

XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.

XX Homo sapiens.
 OS
 XX
 XX WC9856938-A1.
 PN
 XX
 PD 17-DEC-1998.

XX 10-JUN-1998; 98WO-US011927.
 XX 13-JUN-1997; 97US-00874807.
 PR 14-MAY-1998; 98US-00079030.

XX (BAYU) BAYLOR COLLEGE MEDICINE.

PI Guevara JG, Hoogveen RC, Moore JP;
 XX WPI; 1999-070331/06.

XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.

XX Claim 16; Fig 12C; 293pp; English.

XX AAW96827-77 represent nucleic acid binding domains derived from human
 CC apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein component
 CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
 CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
 CC sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis

XX Sequence 36 AA;

Query Match 84.0%; Score 42; DB 2; Length 36;
 Best Local Similarity 90.0%; Pred. No. 0.82;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRLK 10
 | | | | |

DB 11 TRLTRKRLK 20

RESULT 12
 AAW64587
 ID AAW64587 standard; peptide; 37 AA.

XX AAW64587;

XX 23-OCT-1998 (first entry)

XX Human apolipoprotein peptide fragment #1.

XX Factor V; human; detection; protein function; blood coagulation; apo;
 KW fat metabolism; Leyden mutation; deep vein thrombosis; apolipoprotein;
 KW Alzheimer's disease; 5,10-methylenetetrahydrofolate reductase; prion;
 KW hypercystinaemia; factor VII; cardiovascular disease; pathogen; virus.

XX Homo sapiens.

XX EP857973-A2.

XX 12-AUG-1998.

XX 12-JAN-1998; 98EP-00890007.

XX 13-JAN-1997; 97AT-00000044.

XX (IMMO) IMMUNO AG.

XX Moritz B, Kiessig S, Lang H, Schenk V;

XX WPI; 1998-416142/36.

XX Detecting or quantifying mutant protein in presence of wild-type protein
 PT comprises reaction with ligand - used to detect mutant blood coagulation
 PT factors or apolipoproteins for diagnosing risk of thrombosis.

XX Example 2; Page 9; 18pp; German.

XX AAW64587 and AAW64588 are fragments of human apolipoprotein which are
 CC used with Factor V protein fragments in a novel method to detect the
 CC presence of a mutated protein in a sample that may also contain the
 CC corresponding wild-type protein. The method is used to detect mutations
 CC that alter protein functions (either point mutation or small insertions
 CC or deletions), particularly in proteins involved in blood coagulation or
 CC metabolism of fat. Protein functions which are specially detectable are
 CC the Leyden mutation in factor V (associated with increased risk of deep
 CC vein thrombosis), mutations in apolipoprotein (apo) genes (certain
 CC alleles of apoE indicates increased risk of developing Alzheimer's
 CC disease), thermostable 5,10-methylenetetrahydrofolate reductase
 CC (associated with hypercystinaemia and venous thrombosis) and factor VII
 CC mutations (associated with increased risk of cardiovascular disease). The
 CC method can also be applied to proteins from pathogens, e.g. viruses or
 CC prions. The method does not require complex apparatus for polymerase
 CC chain reactions, it is simple, standardisable and reliable and is
 CC particularly suited to routine screening. It also allows mutant protein
 CC in a sample to be quantified

XX Sequence 37 AA;

Query Match 84.0%; Score 42; DB 2; Length 37;
 Best Local Similarity 90.0%; Pred. No. 0.85;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRLK 10
 | | | | |

DB 11 TRLTRKRLK 20

RESULT 13
 AAW6845
 ID AAW6845 standard; peptide; 51 AA.

XX AAW96845;
AC
XX
DT 22-APR-1999 (first entry)
XX
XX Nucleic acid binding domain from apoB-100.
DE
XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
XX
OS Homo sapiens.
XX
XX WO9856938-A1.
PN
XX
XX 17-DEC-1998.
PD
XX
XX 10-JUN-1998; 98WO-US011927.
PF
XX
XX 13-JUN-1997; 97US-00874807.
PR
XX 14-MAY-1998; 98US-00079030.
PR
XX (BAYU) BAYLOR COLLEGE MEDICINE.
PA
XX
XX Guevara JG, Hoogveen RC, Moore JP;
PI WPI; 1999-070331/06.
XX
XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
PT used for delivering nucleic acid to cells for gene therapy and antisense
PT treatment.
XX
XX Claim 16; Page 151; 293pp; English.
PS
XX
XX AAW96827-77 represent nucleic acid binding domains derived from human
CC apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein component
CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
CC sequence can be used in the composition of the invention. The
CC specification describes a composition that comprises LDL and
CC apolipoproteins for the binding and in vivo transport of nucleic acids.
CC The composition is used to deliver nucleic acids to eukaryotic cells, in
CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
CC molecule (or ribozyme). Specifically they are used for gene therapy of
CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
CC fibrosis and arteriosclerosis
XX
SQ Sequence 51 AA;
Query Match 84.0%; Score 42; DB 2; Length 51;
Best Local Similarity 90.0%; Pred. No. 1.2;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TDLTRKRGK 10
Db 6 TRLTRKRGK 15
RESULT 14
ABB37687
ID ABB37687 standard; peptide; 343 AA.
XX
XX ABB37687;
AC
XX
XX 04-FEB-2002 (first entry)
DT
XX
XX Peptide #5193 encoded by human foetal liver single exon probe.
DE
XX
XX Human; foetal liver; gene expression; single exon nucleic acid probe.
KW
XX
XX Homo sapiens.
OS
XX

PN WO200157277-A2.
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US000669.
PF
XX
XX 04-FEB-2000; 2000US-0180312P.
PR
XX 26-MAY-2000; 2000US-0207456P.
PR
XX 30-JUN-2000; 2000US-00608408.
PR
XX 03-AUG-2000; 2000US-00632366.
PR
XX 21-SEP-2000; 2000US-0234687P.
PR
XX 27-SEP-2000; 2000US-0236359P.
PR
XX 04-OCT-2000; 2000GB-00024263.
XX
XX (MOLE-) MOLECULAR DYNAMICS INC.
PA
XX
XX Penn SG, Hanzel DK, Chen W, Rank DR;
PI
XX
XX WPI; 2001-483447/52.
DR
XX
XX Human genome-derived single exon nucleic acid probes useful for analyzing
PT gene expression in human fetal liver.
PT
XX
XX Claim 27; SEQ ID NO 30322; 639pp + Sequence Listing; English.
PS
XX
XX The invention relates to a single exon nucleic acid probe for measuring
CC human gene expression in a sample derived from human foetal liver. The
CC single exon nucleic acid probes may be used for predicting, measuring and
CC displaying gene expression in samples derived from human fetal liver. The
CC present sequence is a peptide encoded by a single exon nucleic acid probe
CC of the invention. Note: The sequence data for this patent did not form
CC part of the printed specification, but was obtained in electronic format
CC directly from WIPO at ftp.wipo.int/pub/published_pct_sequences
XX
SQ Sequence 343 AA;
Query Match 84.0%; Score 42; DB 4; Length 343;
Best Local Similarity 90.0%; Pred. No. 7.8;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TDLTRKRGK 10
Db 169 TRLTRKRGK 178
RESULT 15
ABG52504
ID ABG52504 standard; peptide; 343 AA.
XX
XX ABG52504;
AC
XX
XX 25-FEB-2003 (first entry)
DT
XX
XX Human liver peptide, SEQ ID No 31152.
DE
XX
XX Human; liver; cirrhosis; hyperlipoproteinaemia; hyperlipidaemia;
KW hypercholesterolaemia; coronary heart disease.
KW
XX
XX Homo sapiens.
OS
XX
XX WO200157273-A2.
PN
XX
XX 09-AUG-2001.
PD
XX
XX 30-JAN-2001; 2001WO-US000664.
PF
XX
XX 04-FEB-2000; 2000US-0180312P.
PR
XX 26-MAY-2000; 2000US-0207456P.
PR
XX 30-JUN-2000; 2000US-00608408.
PR
XX 03-AUG-2000; 2000US-00632366.
PR
XX 21-SEP-2000; 2000US-0234687P.
PR
XX 27-SEP-2000; 2000US-0236359P.
PR
XX 04-OCT-2000; 2000GB-00024263.
PR

XX (MOLE-) MOLECULAR DYNAMICS INC.
XX Penn SG, Hanzel DK, Chen W, Rank DR;
XX WPI; 2001-488898/53.
XX Human genome-derived single exon nucleic acid probes useful for analyzing
XX gene expression in human adult liver.
XX Claim 27; SEQ ID NO 31152; 658pp; English.
XX The invention relates to a single exon nucleic acid probe (SENP) (I) for
XX measuring human gene expression in a sample derived from human adult
XX liver, comprising one of 13109 defined nucleotide sequences given in the
XX specification (or complements/ fragments). The probe hybridizes at high
XX stringency to a nucleic acid molecule expressed in the human adult liver.
XX (I) may be used for predicting, measuring and displaying gene expression
XX in samples derived from human adult liver. The genes identified may be
XX involved in genetic liver diseases such as cirrhosis,
XX hyperlipoproteinaemia, hyperlipidaemia and hypercholesterolaemia which is
XX associated with coronary heart disease. ABG47348-ABG59930 represent human
XX liver single exon encoded peptides of the invention. Note: The sequence
XX information for this patent does not appear in the printed specification
XX but was obtained in electronic format directly from WIPO at
XX ftp.wipo.int/pub/published_pct_sequences
XX Sequence 343 AA;

Query Match 84.0%; Score 42; DB 4; Length 343;
Best Local Similarity 90.0%; Pred. No. 7.8;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
Qy 1 TDLTRKRGK 10
Db 169 TDLTRKRGK 178

Search completed: January 13, 2005, 01:43:02
Job time : 78.3798 secs

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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:30:05 ; Search time 14.4262 Seconds
(without alignments)
66.696 Million cell updates/sec

Title: US-09-823-418-12
Perfect score: 50
Sequence: 1 TDLTRKRLK 10
Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues
Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR_79:*
1: pir1:*
2: pir2:*
3: pir3:*
4: pir4:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|----------|--------------------|
| 1 | 44 | 88.0 | 275 | 2 E60950 | apolipoprotein B-1 |
| 2 | 42 | 84.0 | 596 | 2 S32802 | apolipoprotein B - |
| 3 | 42 | 84.0 | 4563 | 1 LPHUB | apolipoprotein B-1 |
| 4 | 40 | 80.0 | 274 | 2 A60950 | apolipoprotein B-1 |
| 5 | 39 | 78.0 | 269 | 2 C60950 | apolipoprotein B-1 |
| 6 | 39 | 78.0 | 779 | 2 JH0102 | apolipoprotein B - |
| 7 | 37 | 74.0 | 1091 | 2 T35822 | probable regulator |
| 8 | 36 | 72.0 | 266 | 2 S60674 | hypothetical prote |
| 9 | 35 | 70.0 | 187 | 2 T21671 | hypothetical prote |
| 10 | 35 | 70.0 | 309 | 1 E65112 | hypothetical 34.6 |
| 11 | 35 | 70.0 | 309 | 2 E85985 | hypothetical prote |
| 12 | 35 | 70.0 | 309 | 2 B91140 | hypothetical prote |
| 13 | 35 | 70.0 | 382 | 1 A44056 | nucleoside prote |
| 14 | 35 | 70.0 | 1217 | 2 T25894 | hypothetical prote |
| 15 | 35 | 70.0 | 3450 | 2 T26963 | hypothetical prote |
| 16 | 35 | 70.0 | 3461 | 2 T26964 | hypothetical prote |
| 17 | 34 | 68.0 | 168 | 2 T03168 | hypothetical prote |
| 18 | 34 | 68.0 | 206 | 2 T01788 | aminoglycoside 6'- |
| 19 | 34 | 68.0 | 211 | 2 F75474 | hypothetical prote |
| 20 | 34 | 68.0 | 493 | 2 E71008 | hypothetical prote |
| 21 | 34 | 68.0 | 680 | 2 S29682 | DNA topoisomerase |
| 22 | 34 | 68.0 | 684 | 2 S29683 | DNA gyrase B, novo |
| 23 | 34 | 68.0 | 686 | 2 T10969 | DNA topoisomerase |
| 24 | 34 | 68.0 | 1058 | 2 S65460 | apolipoprotein B - |
| 25 | 34 | 68.0 | 1253 | 1 A44400 | myosin heavy chain |
| 26 | 34 | 68.0 | 1254 | 2 A54818 | myosin-VI [similar |
| 27 | 34 | 68.0 | 1265 | 2 A59299 | unconventional myo |
| 28 | 34 | 68.0 | 1615 | 2 JC6510 | ras-responsive ele |
| 29 | 34 | 68.0 | 1778 | 2 JT0382 | apolipoprotein B - |

| | | | | | |
|----|----|------|------|----------|--------------------|
| 30 | 34 | 68.0 | 2629 | 2 I46569 | apolipoprotein B - |
| 31 | 33 | 66.0 | 102 | 2 D75018 | hypothetical prote |
| 32 | 33 | 66.0 | 254 | 2 H95070 | hypothetical prote |
| 33 | 33 | 66.0 | 264 | 2 F97938 | hypothetical prote |
| 34 | 33 | 66.0 | 325 | 2 B72475 | probable transcrip |
| 35 | 33 | 66.0 | 339 | 2 S62596 | ubiquinol-cytochro |
| 36 | 33 | 66.0 | 391 | 2 S60672 | replication protei |
| 37 | 33 | 66.0 | 402 | 2 S55980 | probable membrane |
| 38 | 33 | 66.0 | 784 | 2 JH0101 | apolipoprotein B-1 |
| 39 | 33 | 66.0 | 2100 | 2 T03223 | probable polyketid |
| 40 | 32 | 64.0 | 70 | 2 B64497 | hypothetical prote |
| 41 | 32 | 64.0 | 149 | 2 E72338 | conserved hypotnet |
| 42 | 32 | 64.0 | 214 | 2 AE2934 | transcription regu |
| 43 | 32 | 64.0 | 219 | 2 B98348 | probable transcrip |
| 44 | 32 | 64.0 | 252 | 2 AE0876 | conserved hypotnet |
| 45 | 32 | 64.0 | 281 | 2 F82832 | pantoate-beta-alan |

ALIGNMENTS

RESULT 1

E60950
apolipoprotein B-100 - chicken (fragment)
C/Species: Gallus gallus (chicken)
C/Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C/Accession: E60950
R/Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A/Title: A cross-species comparison of the apolipoprotein B domain that binds to the Lp
A/Reference number: A60950; MUID:90324804; PMID:2373961
A/Accession: E60950
A/Molecule type: mRNA
A/Residues: 1-275 <LAN>
A/Cross-references: UNIPROT:Q7LZ77
C/Superfamily: apolipoprotein B
C/Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 88.0%; Score 44; DB 2; Length 275;
Best Local Similarity 90.0%; Pred. No. 0.46; 1; Indels 0; Gaps 0;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLK 10
| | | | | | | | | |
Db 221 TSLTRKRLK 230

RESULT 2

S32802
apolipoprotein B - crab-eating macaque (fragment)
C/Species: Macaca fascicularis (crab-eating macaque)
C/Date: 06-Jan-1995 #sequence_revision 08-Jan-1995 #text_change 09-Jul-2004
C/Accession: S32802
R/Pape, M.E.; Castle, C.K.; Murray, R.W.; Funk, G.M.; Hunt, C.E.; Marotti, K.R.; Melchior
Biochim. Biophys. Acta 1086, 326-334, 1991
A/Title: Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional
A/Reference number: S32802; MUID:92075708; PMID:1742325
A/Accession: S32802
A/Status: preliminary
A/Molecule type: mRNA
A/Residues: 1-596 <PAP>
A/Cross-references: UNIPROT:Q28473; EMBL:X15737; NID:g38047; PIDN:CAA33755.1; PID:g9301

Query Match 84.0%; Score 42; DB 2; Length 596;
Best Local Similarity 90.0%; Pred. No. 2.4; 1; Indels 0; Gaps 0;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLK 10
| | | | | | | | | |
Db 226 TSLTRKRLK 235

RESULT 3

LPHUB
apolipoprotein B-100 precursor - human
N:Contains: apolipoprotein B-26; apolipoprotein B-48; apolipoprotein B-74
C:Species: Homo sapiens (man)
C>Date: 28-Dec-1987 #sequence_revision 28-Dec-1987 #text_change 09-Jul-2004
A:Accession: A25679; A25263; A25267; A25266; A24320; A24684; A23817; A25774; A264452; I61909; I79510; I39474; I39469; I84624; I37179; P80058
R:Ludwig, E.H.; Blackhart, B.D.; Pierotti, V.R.; Caiati, L.; Fortier, C.; Knott, T.; Scott DNA 6, 363-372, 1987
A:Title: DNA sequence of the human apolipoprotein B gene.
A:Reference number: A27850; MUID:88003974; PMID:3652907
A:Accession: A27850
A:Molecule type: DNA
A:Residues: 1-617, 'A', 619-1929, 'F', 1931-3318, 'D', 3320-3426, 'T', 3428-3431, 'Q', 3433-3731, 'A'
A:Cross-references: UNIPROT:P04114; UNIPROT:P78482; UNIPROT:P78479; UNIPROT:Q9UNN0; UNIPROT:Q9UNN1
R:Cladaras, C.; Hadzopoulou-Cladaras, M.; Nolte, R.T.; Atkinson, D.; Zannis, V.I. EMBO J. 5, 3495-3507, 1986
A:Title: The complete sequence and structural analysis of human apolipoprotein B-100: re
A:Reference number: A91058; MUID:87161758; PMID:3030729
A:Accession: A25679
A:Molecule type: mRNA
A:Residues: 1-11, 15-2539, 'S', 2541-3823, 'R', 3825-4563 <CLA>
A>Note: 1109-Asp was also found
R:Knott, T.J.; Wallis, S.C.; Powell, L.M.; Pease, R.J.; Lusis, A.J.; Blackhart, B.; McCa Nucleic Acids Res. 14, 7501-7503, 1986
A:Title: Complete cDNA and derived protein sequence of human apolipoprotein B-100.
A:Reference number: A93639; MUID:87016385; PMID:3763409
A:Accession: A25263
A:Molecule type: mRNA
A:Residues: 1-272, 'N', 274-617, 'A', 619-1217, 'E', 1219-2091, 'V', 2093-2364, 'T', 2366-2679, 'Q'
A:Cross-references: GB:X04506; NID:G34330; PIDN:CAA28191.1; PID:G34331
R:Law, S.W.; Grant, S.M.; Higuchi, K.; Hospattankar, A.; Lackner, K.; Lee, N.; Brewer J Proc. Natl. Acad. Sci. U.S.A. 83, 8142-8146, 1986
A:Title: Human liver apolipoprotein B-100 cDNA: complete nucleic acid and derived amino A:Reference number: A94134; MUID:87041416; PMID:3464946
A:Accession: A25267
A:Molecule type: mRNA
A:Residues: 1-617, 'A', 619-703, 'P', 705-792, 'R', 794-1270, 'S', 1272-1866, 'G', 1868-2036, 'N', 24189-4220, 'M', 4222-4563 <LAW>
A>Note: the codons given for residues 704, 793, 1271, 2037, 2933, 3286, 3782, 4188, and R:Chen, S.H.; Yang, C.Y.; Chen, P.F.; Setzer, D.; Tanimura, M.; Li, W.H.; Gotto Jr., A.M J. Biol. Chem. 261, 12918-12921, 1986
A:Title: The complete cDNA and amino acid sequence of human apolipoprotein B-100.
A:Reference number: A92556; MUID:87008488; PMID:3759943
A:Accession: A25266
A:Molecule type: mRNA
A:Residues: 1-97, 'I', 99-328, 'V', 330-644, 'I', 646-918, 'P', 920-3318, 'D', 3320-3426, 'T', 3428-9-4132, 'G', 4134-4180, 'E', 4182-4563 <CHE>
A:Cross-references: GB:J02610; NID:G178803; PIDN:AAA35549.1; PID:G178804
A>Note: a total of 2366 residues were confirmed by direct sequencing of tryptic peptides R:Protter, A.A.; Hardman, D.A.; Sato, K.Y.; Schilling, J.W.; Yamanaka, M.; Hort, Y.J.; H Proc. Natl. Acad. Sci. U.S.A. 83, 5678-5682, 1986
A:Title: Analysis of cDNA clones encoding the entire B-26 region of human apolipoprotein A:Reference number: A24320; MUID:86287319; PMID:3461454
A:Accession: A24320
A:Molecule type: mRNA
A:Residues: 1-97, 'I', 99-617, 'A', 619-941, 'YYIWSLPKP', 951-1138, 'PTGRLPNCFSNGLICYSLWHSQOE A:Cross-references: GB:M14081; NID:G178795; PIDN:AAA51752.1; PID:G553189
R:Law, S.W.; Lackner, K.J.; Hospattankar, A.V.; Anchors, J.M.; Sakaguchi, A.Y.; Naylor, Proc. Natl. Acad. Sci. U.S.A. 82, 8340-8344, 1985
A:Title: Human apolipoprotein B-100: cloning, analysis of liver mRNA, and assignment of A:Reference number: A24684; MUID:86094221; PMID:3001697
A:Accession: A24684
A:Molecule type: mRNA
A:Residues: 485-617, 'A', 619-1044 <LAW>
A:Cross-references: GB:M12480; NID:G178791; PIDN:AAA51751.1; PID:G178792
R:Protter, A.A.; Hardman, D.A.; Schilling, J.W.; Miller, J.; Appleby, V.; Chen, G.C.; K Proc. Natl. Acad. Sci. U.S.A. 83, 1467-1471, 1986
A:Title: Isolation of a cDNA clone encoding the amino-terminal region of human apolipop A:Reference number: A94088; MUID:86149325; PMID:3513177
A:Accession: A23817
A:Molecule type: mRNA

A;Residues: 1-291 <PRO>
A;Cross-references: GB:M12681; NID:G178797; PIDN:AAA51753.1; PID:G178798
R;Deeb, S.S.; Motulsky, A.G.; Albers, J.J. Proc. Natl. Acad. Sci. U.S.A. 82, 4983-4986, 1985
A;Title: A partial cDNA clone for human apolipoprotein B.
A;Reference number: A25774; MUID:85270450; PMID:3860836
A;Accession: A25774
A:Molecule type: mRNA
A;Residues: 709-791, 'SSSWKAASHGCPHSAGD', 810-906 <DEE>
A;Cross-references: GB:K03175; NID:G178821; PIDN:AAA51759.1; PID:G178822
R;Carlsson, P.; Darnfors, C.; Olofsson, S.O.; Bjursell, G. Gene 49, 29-51, 1986
A;Title: Analysis of the human apolipoprotein B gene; complete structure of the B-74 re A;Reference number: A91565; MUID:87191999; PMID:2883086
A;Accession: A26533
A:Molecule type: mRNA
A;Residues: 1282-2721, 2742-3280, 'L', 3292-3336, 'N', 3338-3948, 'F', 3950-3963, 'Y', 3965-4180, 'A'
A:Cross-references: GB:M15421; NID:G178817; PIDN:AAA51758.1; PID:G178818
R;Hardman, D.A.; Protter, A.A.; Chen, G.C.; Schilling, J.W.; Sato, K.Y.; Lau, K.; Yaman Biochemistry 26, 5478-5486, 1987
A;Title: Structural comparison of human apolipoproteins B-48 and B-100.
A;Reference number: A23671; MUID:88050832; PMID:3676265
A;Accession: A23671
A:Molecule type: mRNA
A;Residues: 1671-2323, 'PYW', 2327-2352, 'H', 2354-2398 <HAR>
A;Cross-references: GB:M17367; NID:G178731; PIDN:AAA51741.1; PID:G178732
R;Shoulders, C.C.; Myant, N.B.; Sidoli, A.; Rodriguez, J.C.; Cortese, C.; Baralle, P.E. Atherosclerosis 58, 277-289, 1985
A;Title: Molecular cloning of human LDL apolipoprotein B cDNA. Evidence for more than o A;Reference number: A90084; MUID:86130855; PMID:3841481
A;Accession: A29287
A:Molecule type: mRNA
A;Residues: 3846-4298 <SHO>
R;Pfizner, R.; Wagener, R.; Stoffel, W. Biol. Chem. Hoppe-Seyler 367, 1077-1083, 1986
A;Title: Isolation, expression and characterization of a human apolipoprotein B 100-spe A;Reference number: A25572; MUID:87076044; PMID:3024665
A;Accession: A25572
A:Molecule type: mRNA
A;Residues: 4219-4337, 'S', 4339-4563 <PFI>
A;Cross-references: GB:M36676
R;Wei, C.F.; Chen, S.H.; Yang, C.Y.; Marcel, Y.L.; Milne, R.W.; Li, W.H.; Sparrow, J.T. Proc. Natl. Acad. Sci. U.S.A. 82, 7265-7269, 1985
A;Reference number: A24738; MUID:86042646; PMID:2932736
A;Accession: A24738
A:Molecule type: mRNA
A;Residues: 'N', 3729-3731, 'I', 3733-3875, 'A', 3877-3948, 'F', 3950-3963, 'Y', 3965-3982, 'S', 3 R;Chen, S.H.; Habib, G.; Yang, C.Y.; Gu, Z.W.; Lee, B.R.; Weng, S.; Silberman, S.R.; Ca Science 238, 363-366, 1987
A;Title: Apolipoprotein B-48 is the product of a messenger RNA with an organ-specific i A;Reference number: A40133; MUID:88018019; PMID:3659919
A;Accession: B40133
A:Molecule type: mRNA
A;Residues: 2165-2179 <CHI>
A;Cross-references: GB:M18036; NID:G178799; PIDN:AAA51754.1; PID:G178800
A>Note: this mRNA includes the stop codon of the organ-specific mRNA for apo48 A;Accession: A40133
A:Molecule type: protein
A;Residues: 51-75;101-110;129-139;158-174;197-207;276-287;298-304;306-314;526-532;538-5 36;1486-1498;1537-1556;1563-1572;1601-1610;1647-1661;1697-1724;1770-1781;1859-1897;1968
A>Note: these fragments were derived from apo48
R;Hardman, D.A.; Protter, A.A.; Schilling, J.W.; Kane, J.P. Biochem. Biophys. Res. Commun. 149, 1214-1219, 1987
A;Title: Carboxyl terminal analysis of human B-48 protein confirms the novel mechanism A;Reference number: A28002; MUID:88106542; PMID:3426612
A;Accession: A28002
A:Molecule type: mRNA
A;Residues: 2129-2179, 2181-2235 <HA>
A;Cross-references: GB:M18471
A;Experimental source: intestine
A>Note: this mRNA from intestine includes a stop codon created by RNA editing in place , R;Mehrabian, M.; Schumaker, V.N.; Fareed, G.C.; West, R.; Johnson, D.F.; Kirchgessner,

Nucleic Acids Res. 13, 6937-6953, 1985
 A>Title: Human apolipoprotein B: identification of cDNA clones and characterization of m
 A:Reference number: A24269; MUID:86041888; PMID:3903660
 A:Accession: A24269
 A:Molecule type: mRNA
 A:Residues: 3056-3159 <MEH>
 A:Cross-references: GB:X03045; NID:928783; PIDN:CAA26850.1; PID:9292609
 R:Hospatankar, A.V.; Higuchi, K.; Law, S.W.; Meglin, N.; Brewer Jr., H.B.
 Biochem. Biophys. Res. Commun. 148, 279-285, 1987
 A>Title: Identification of a novel in-frame translational stop codon in human intestine
 A:Reference number: A29659; MUID:88049670; PMID:2445342
 A:Accession: A29659
 A:Molecule type: mRNA
 A:Residues: 2169-2179 <HOS>
 A>Note: the sequence shown represents the carboxyl end of apolipoprotein B-48
 A>Note: two RNA species, 14.1kb and 7.5kb in length, were isolated from the human intest
 ch encodes the 250K apoB-48, CAA encoding 2180-Gln is substituted by the stop codon TAA,
 R:Yang, C.; Kim, T.W.; Wang, S.; Lee, B.; Yang, M.; Goto Jr., A.M.
 Proc. Natl. Acad. Sci. U.S.A. 87, 5523-5527, 1990
 A>Title: Isolation and characterization of sulfhydryl and disulfide peptides of human ap
 A:Reference number: A35783; MUID:90319144; PMID:2115173
 A:Contents: disulfide bonds
 A:Accession: A35783
 A:Molecule type: protein
 A:Residues: 28-41;76-97, I', 99-100;175-193;206-215;239-249;259-266;357-399;455-490;512-5
 A>Note: cysteines at positions 1112, 1422, 1505, 1662, 3761, 3917, and 4217 have free su
 R:LeBoeuf, R.C.; Miller, C.; Shively, J.E.; Schumaker, V.N.; Balla, M.A.; Lusis, A.J.
 FEBS Lett. 170, 105-108, 1984
 A>Title: Human apolipoprotein B: partial amino acid sequence.
 A:Reference number: A22006; MUID:84208786; PMID:6373369
 A:Accession: A22006
 A:Molecule type: protein
 A:Residues: 873-892, 'K', 894-896 <LE1>
 A:Accession: B22006
 A:Molecule type: protein
 A:Residues: 3113, 'L', 3115-3130, 'R', 3132-3133, 'P', 3135-3136, 'R' <LE2>
 R:Blackhart, B.D.; Ludwig, E.M.; Pierotti, V.R.; Calati, L.; Onasch, M.A.; Wallis, S.C.;
 J. Biol. Chem. 261, 15364-15367, 1986
 A>Title: Structure of the human apolipoprotein B gene.
 A:Reference number: A92564; MUID:87057153; PMID:2945672
 A:Contents: annotation; gene structure
 R:Wagener, R.; Pflitzner, R.; Stoffel, W.
 Biol. Chem. Hoppe-Seyler 368, 419-425, 1987
 A>Title: Studies on the organization of the human apolipoprotein B 100 gene.
 A:Reference number: A90715; MUID:87271140; PMID:2886136
 A:Contents: annotation; gene structure
 R:Weisgraber, K.H.; Rall Jr., S.C.
 J. Biol. Chem. 262, 11097-11103, 1987
 A>Title: Human apolipoprotein B-100 heparin-binding sites.
 A:Reference number: A92605; MUID:87280197; PMID:3301850
 A:Contents: annotation; heparin binding and disulfide bond
 R:Dashti, N.; Lee, D.M.; Mok, T.
 Biochem. Biophys. Res. Commun. 137, 493-499, 1986
 A>Title: Apolipoprotein B is a calcium binding protein.
 A:Reference number: A30125; MUID:86242245; PMID:3087360
 A:Contents: annotation; calcium binding
 R:Carlsson, P.; Olofsson, S.O.; Bondjers, G.; Darnfors, C.; Wiklund, O.; Bjursell, G.
 Nucleic Acids Res. 13, 8813-8825, 1985
 A>Title: Molecular cloning of human apolipoprotein B cDNA.
 A:Reference number: 137178; MUID:86093680; PMID:3841204
 A:Accession: 137180

Query Match 84.0%; Score 42; DB 1; Length 4563;
 Best Local Similarity 90.0%; Pred. No. 16;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLGLK 10

|||||

Db 3385 TRLTRKRLGLK 3394

RESULT 4

A60950
 apolipoprotein B-100 - rabbit (fragment)
 C:Species: Oryctolagus cuniculus (domestic rabbit)
 C>Date: 31-Dec-1993 #sequence_revision 09-Sep-1994 #text_change 09-Jul-2004
 C:Accession: A60950
 R:Law, A.; Scott, J.
 J. Lipid Res. 31, 1109-1120, 1990
 A>Title: A cross-species comparison of the apolipoprotein B domain that binds to the Lf
 A:Reference number: A60950; MUID:90324804; PMID:2373961
 A:Accession: A60950
 A:Molecule type: mRNA
 A:Residues: 1-274 <LAW>
 A:Cross-references: UNIPROT:Q7M2U9
 A>Note: authors translated the codon GAT for residue 155 as His
 C:Superfamily: apolipoprotein B
 C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 80.0%; Score 40; DB 2; Length 274;
 Best Local Similarity 80.0%; Pred. No. 2.9;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLGLK 10

|||||

Db 221 SSLTRKRLGLK 230

RESULT 5

C60950
 apolipoprotein B-100 - golden hamster (fragment)
 C:Species: Mesocricetus auratus (golden hamster)
 C>Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
 C:Accession: C60950
 R:Law, A.; Scott, J.
 J. Lipid Res. 31, 1109-1120, 1990
 A>Title: A cross-species comparison of the apolipoprotein B domain that binds to the Lf
 A:Reference number: A60950; MUID:90324804; PMID:2373961
 A:Accession: C60950
 A:Molecule type: DNA
 A:Residues: 1-269 <LAW>
 A:Cross-references: UNIPROT:Q60537; UNIPROT:Q60536
 C:Superfamily: apolipoprotein B
 C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein

Query Match 78.0%; Score 39; DB 2; Length 269;
 Best Local Similarity 100.0%; Pred. No. 4.5;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LTRKRLGLK 10

|||||

Db 218 LTRKRLGLK 225

RESULT 6

JH0102
 apolipoprotein B - golden hamster (fragment)
 C:Species: Mesocricetus auratus (golden hamster)
 C>Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
 C:Accession: JH0102
 R:Smith, T.J.
 submitted to GenBank, June 1990
 A:Reference number: A38864
 A:Accession: JH0102
 A:Molecule type: DNA
 A:Residues: 1-779 <SMI>
 A:Cross-references: UNIPROT:Q60536; GB:M35187
 A>Note: this is a revision to the sequence from reference JH0101
 R:Smith, T.J.; Hautamaa, D.; Maeda, N.
 Gene 87, 309-310, 1990

Query Match 84.0%; Score 42; DB 1; Length 4563;
 Best Local Similarity 90.0%; Pred. No. 16;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRLGLK 10

|||||

Db 3385 TRLTRKRLGLK 3394

RESULT 4

A:Contents: annotation
 A>Note: this sequence has been revised in reference A38864
 C:Genetics:

A:Gene: apoB
 C:Superfamily: apolipoprotein B
 C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;
 F:435-445/Region: receptor binding
 F:646-656/Region: receptor binding

Query Match 78.0%; Score 39; DB 2; Length 779;
 Best Local Similarity 100.0%; Pred. No. 12;
 Matches 8; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 LTRKRGGLK 10
 |||||
 Db 644 LTRKRGGLK 651

RESULT 7

T35822
 probable regulatory protein - Streptomyces coelicolor
 C:Species: Streptomyces coelicolor
 C:Date: 05-Nov-1999 #sequence_revision 05-Nov-1999 #text_change 09-Jul-2004
 C:Accession: T35822
 R:Murphy, L.; Harris, D.; Bentley, S.D.; Parkhill, J.; Barrell, B.G.; Rajandream, M.A.
 submitted to the EMBL Data Library, February 1999
 A:Reference number: Z21589
 A:Accession: T35822
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-1091 <MUR>
 A:Cross-references: UNIPROT:Q92573; EMBL:AL035569; PIDN:CA837582.1; GSPDB:GN000070; SCORE
 A:Experimental source: strain A3(2)
 C:Genetics:
 A:Gene: SCOEDB:SC8D9.18

Query Match 74.0%; Score 37; DB 2; Length 1091;
 Best Local Similarity 87.5%; Pred. No. 41;
 Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 2 DLTRKRGL 9
 |||||
 Db 729 DLTRRGL 736

RESULT 8

S60674
 hypothetical protein B - Corynebacterium glutamicum plasmid pGA1
 C:Species: Corynebacterium glutamicum
 C:Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 09-Jul-2004
 C:Accession: S60674
 R:Nesvera, J.; Patek, M.; Hochmannova, J.; Abrahamova, Z.
 submitted to the EMBL Data Library, August 1995
 A:Description: Complete nucleotide sequence of the cryptic plasmid pGA1 from Corynebacte
 A:Reference number: S60673
 A:Accession: S60674
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-266 <NES>
 A:Cross-references: UNIPROT:Q46059; EMBL:X90817; NID:9951006; PIDN:CAA62329.1; PID:99510
 C:Genetics:
 A:Genome: plasmid pGA1

Query Match 72.0%; Score 36; DB 2; Length 266;
 Best Local Similarity 66.7%; Pred. No. 18;
 Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

QY 2 DLTRKRGLK 10
 |||||
 Db 211 DLGRKKGIK 219

RESULT 9

T21671
 hypothetical protein F32H2.6 - Caenorhabditis elegans
 C:Species: Caenorhabditis elegans

C:Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
 C:Accession: T21671
 R:Kershaw, J.
 submitted to the EMBL Data Library, November 1996
 A:Reference number: Z19457
 A:Accession: T21671
 A:Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-187 <WIL>
 A:Cross-references: UNIPROT:P91866; EMBL:Z81523; PIDN:CA804239.1; GSPDB:GN000019; CESP:F
 A:Experimental source: clone F32H2
 C:Genetics:
 A:Gene: CESP:F32H2.6
 A:Map position: 1
 A:Introns: 22/2; 103/1

Query Match 70.0%; Score 35; DB 2; Length 187;
 Best Local Similarity 70.0%; Pred. No. 20;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
 |||||
 Db 54 TDLPKRGRKK 63

RESULT 10

E65112
 hypothetical 34.6 kD protein in arcB-gltB intergenic region - Escherichia coli (strain :
 C:Species: Escherichia coli
 C:Date: 10-Sep-1999 #sequence_revision 10-Sep-1999 #text_change 09-Jul-2004
 C:Accession: E65112
 R:Blattner, F.R.; Plunkett III, G.; Bloch, C.A.; Perna, N.T.; Burland, V.; Riley, M.; C;
 A:Rose, D.J.; Mau, B.; Shao, Y.
 Science 277, 1453-1462, 1997
 A:Title: The complete genome sequence of Escherichia coli K-12.
 A:Reference number: A64720; MUID:97426617; PMID:9278503
 A:Accession: E65112
 A:Status: preliminary; nucleic acid sequence not shown; translation not shown
 A:Molecule type: DNA
 A:Residues: 1-309 <BLAT>
 A:Cross-references: UNIPROT:P45476; GB:AE000400; GB:U00096; NID:92367203; PIDN:AAC76243
 A:Experimental source: strain K-12, substrain MG1655
 C:Genetics:
 A:Gene: yhcC
 C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 70.0%; Score 35; DB 1; Length 309;
 Best Local Similarity 70.0%; Pred. No. 32;
 Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
 |||||
 Db 170 TQLARQGLK 179

RESULT 11

E85985
 hypothetical protein yhcC [imported] - Escherichia coli (strain O157:H7, substrain EDL9
 C:Species: Escherichia coli
 C:Date: 16-Feb-2001 #sequence_revision 16-Feb-2001 #text_change 09-Jul-2004
 C:Accession: E85985
 R:Perna, N.T.; Plunkett III, G.; Burland, V.; Mau, B.; Glaesner, J.D.; Rose, D.J.; Mayhe
 iller, L.; Grobeck, E.J.; Davis, N.W.; Lim, A.; Dinalanta, E.; Potamouisis, K.; Apodaca
 Nature 409, 529-533, 2001
 A:Title: Genome sequence of enterohemorrhagic Escherichia coli O157:H7.
 A:Reference number: AB5480; MUID:21074935; PMID:11206551
 A:Accession: E85985
 A:Status: preliminary
 A:Molecule type: DNA
 A:Residues: 1-309 <STO>
 A:Cross-references: UNIPROT:P45476; GB:AE005174; NID:912517832; PIDN:AA058345.1; GSPDB:B
 A:Experimental source: strain O157:H7, substrain EDL933
 C:Genetics:

A:Gene: yhcC
C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 70.0%; Score 35; DB 2; Length 309;
Best Local Similarity 70.0%; Pred. No. 32;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
| | | | |
DB 170 TQLARQRLGK 179

RESULT 12
B91140
hypothetical protein ECs4090 [imported] - Escherichia coli (strain O157:H7, substrain R1
C:Species: Escherichia coli
C>Date: 18-Jul-2001 #sequence_revision 18-Jul-2001 #text_change 09-Jul-2004
C:Accession: B91140
R:Hayashi, T.; Makino, K.; Ohnishi, M.; Kurokawa, K.; Ishii, K.; Yokoyama, K.; Han, C.G.
gasawara, N.; Yasunaga, T.; Kuhara, S.; Shiba, T.; Hattori, M.; Shinagawa, H.
DNA Res. 8, 11-22, 2001
A:Title: Complete genome sequence of enterohemorrhagic Escherichia coli O157:H7 and gen
A:Reference number: A99629; MUID:21156231; PMID:11258796
A:Accession: B91140
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-309 <HAY>
A:Cross-references: UNIPROT:P45476; GB:BA000007; PIDN:BA037513.1; PID:gl3363563; GSPDB:C
A:Experimental source: strain O157:H7, substrain R1MD 0509952
C:Genetics:
A:Gene: ECs4090
C:Superfamily: Methanococcus jannaschii conserved hypothetical protein MJ0486

Query Match 70.0%; Score 35; DB 2; Length 309;
Best Local Similarity 70.0%; Pred. No. 32;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 1 TDLTRKRGGLK 10
| | | | |
DB 170 TQLARQRLGK 179

RESULT 13
A44056
nucleocapsid protein - canine coronavirus (strain K378)
C:Alternate names: N protein
C:Species: canine coronavirus
C>Date: 30-Sep-1993 #sequence_revision 30-Sep-1993 #text_change 09-Jul-2004
C:Accession: A44056
R:Vennema, H.; Rossen, J.W.A.; Wesseling, J.; Horzinek, M.C.; Rottier, P.J.M.
Virology 191, 134-140, 1992
A:Title: Genomic organization and expression of the 3' end of the canine and feline ente
A:Reference number: A44056; MUID:93033103; PMID:1329312
A:Accession: A44056
A:Molecule type: genomic RNA
A:Residues: 1-382 <VEN>
A:Cross-references: UNIPROT:Q04700; GB:X66717; NID:g58849; PIDN:CAA47246.1; PID:g58850
C:Genetics:
A:Gene: N
C:Superfamily: coronavirus nucleocapsid protein
C:Keywords: glycoprotein; nucleocapsid
F:28,134,154,172,364/Binding site: carbohydrate (Asn) (covalent) #status predicted

Query Match 70.0%; Score 35; DB 1; Length 382;
Best Local Similarity 75.0%; Pred. No. 39;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 DLTTRKRG 9
| | | | |
DB 13 DITKRG 20

T25894
hypothetical protein T19B4.2 - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 09-Jul-2004
C:Accession: T25894
R:Gattung, S.
submitted to the EMBL Data Library, November 1996
A:Description: The sequence of C. elegans cosmid T19B4.
A:Reference number: Z20106
A:Accession: T25894
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-1217 <GAT>
A:Cross-references: UNIPROT:P91457; EMBL:U80438; PIDN:AAB37636.1; GSPDB:GN00019; CESP:
A:Experimental source: strain Bristol N2; clone T19B4
C:Genetics:
A:Gene: CESP:T19B4.2
A:Map position: 1
A:Introns: 66/1; 119/3; 321/3; 552/3; 1123/2; 1197/2

Query Match 70.0%; Score 35; DB 2; Length 1217;
Best Local Similarity 100.0%; Pred. No. 1.2e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TDLTRKR 7
| | | | |
DB 290 TDLTRKR 296

RESULT 15
T26963
hypothetical protein ZK1151.2a - Caenorhabditis elegans
C:Species: Caenorhabditis elegans
C>Date: 15-Oct-1999 #sequence_revision 15-Oct-1999 #text_change 29-Oct-1999
C:Accession: T26963; T27704
R:Harris, B.
submitted to the EMBL Data Library, September 1998
A:Reference number: Z20292
A:Accession: T26963
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-3450 <WIL>
A:Cross-references: EMBL:AL031637; PIDN:CAA21049.1; GSPDB:GN00019; CESP:ZK1151.2a
A:Experimental source: clone Y47H9B
R:Harris, B.
submitted to the EMBL Data Library, March 1997
A:Reference number: Z20408
A:Accession: T27704
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-3450 <WIL>
A:Cross-references: EMBL:Z93398; PIDN:CAB07725.1; GSPDB:GN00019; CESP:ZK1151.2a
A:Experimental source: clone ZK1151
C:Genetics:
A:Gene: CESP:ZK1151.2a
A:Map position: 1
A:Introns: 270/3; 757/2; 1105/1; 1259/3; 2312/2; 2613/1; 2825/2; 3180/1; 3217/3; 3257/3

Query Match 70.0%; Score 35; DB 2; Length 3450;
Best Local Similarity 75.0%; Pred. No. 3e+02;
Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 DLTTRKRG 9
| | | | |
DB 2762 DVTTRKRG 2769

Search completed: January 13, 2005, 01:52:39
Job time : 15.4262 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 78.0328 Seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-12
Perfect score: 50
Sequence: 1 TDLTRKRLK 10

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : UniProt_02.*

1: uniprot_sprot.*

2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------------|--------------------|
| 1 | 44 | 88.0 | 275 | 2 Q7LZ77 | Q7LZ77 gallus gall |
| 2 | 44 | 88.0 | 387 | 2 Q7YQN2 | Q7YQN2 phalanger o |
| 3 | 44 | 88.0 | 400 | 2 Q7YQW9 | Q7YQW9 ornithorhyn |
| 4 | 44 | 88.0 | 405 | 2 Q7YQNO | Q7YQNO tachyglossu |
| 5 | 42 | 84.0 | 414 | 2 Q7YQRS | Q7YQRS aotus vocif |
| 6 | 42 | 84.0 | 596 | 2 Q28473 | Q28473 macaca fasc |
| 7 | 42 | 84.0 | 3262 | 2 Q13788 | Q13788 homo sapien |
| 8 | 42 | 84.0 | 4563 | 1 APB_HUMAN | P04114 homo sapien |
| 9 | 42 | 84.0 | 4563 | 2 Q7Z600 | Q7Z600 homo sapien |
| 10 | 40 | 80.0 | 263 | 2 Q7YQO0 | Q7YQO0 procavia ca |
| 11 | 40 | 80.0 | 274 | 2 Q7M2U9 | Q7M2U9 oryctolagus |
| 12 | 40 | 80.0 | 304 | 2 Q7YQF9 | Q7YQF9 echinops te |
| 13 | 40 | 80.0 | 314 | 2 Q7YQW8 | Q7YQW8 ictonyx str |
| 14 | 40 | 80.0 | 316 | 2 Q7YQP3 | Q7YQP3 nandinia bi |
| 15 | 40 | 80.0 | 318 | 2 Q7YQW9 | Q7YQW9 zalophus ca |
| 16 | 40 | 80.0 | 319 | 2 Q7YQO0 | Q7YQO0 vulpes vulp |
| 17 | 40 | 80.0 | 319 | 2 Q7YQF2 | Q7YQF2 panthera le |
| 18 | 40 | 80.0 | 320 | 2 Q7YQP4 | Q7YQP4 manis sp. k |
| 19 | 40 | 80.0 | 322 | 2 Q7YQP5 | Q7YQP5 manis sp. k |
| 20 | 40 | 80.0 | 339 | 2 Q7YR05 | Q7YR05 macroselid |
| 21 | 40 | 80.0 | 361 | 2 Q7YQP8 | Q7YQP8 amblysomus |
| 22 | 40 | 80.0 | 364 | 2 Q7YQO1 | Q7YQO1 dugong dugo |
| 23 | 40 | 80.0 | 386 | 2 Q7YQR1 | Q7YQR1 tupia tana |
| 24 | 40 | 80.0 | 392 | 2 Q7YR11 | Q7YR11 tarsius syr |
| 25 | 40 | 80.0 | 411 | 2 Q7YQP7 | Q7YQP7 ochotona pr |
| 26 | 40 | 80.0 | 422 | 2 Q7YR12 | Q7YR12 talpa europ |
| 27 | 40 | 80.0 | 423 | 2 Q7YQO9 | Q7YQO9 sorex monti |
| 28 | 40 | 80.0 | 426 | 2 Q7YQR2 | Q7YQR2 aices aices |
| 29 | 40 | 80.0 | 429 | 2 Q7YQO8 | Q7YQO8 crocidura f |
| 30 | 40 | 80.0 | 438 | 2 Q7YQR4 | Q7YQR4 balaena mys |
| 31 | 40 | 80.0 | 441 | 2 Q7YQR3 | Q7YQR3 phocoenoide |

| | | | | | |
|----|----|------|-----|----------|---------------------|
| 32 | 40 | 80.0 | 443 | 2 Q7YQNS | Q7YQNS megaderma l |
| 33 | 40 | 80.0 | 443 | 2 Q7YQP6 | Q7YQP6 lepus ameri |
| 34 | 40 | 80.0 | 445 | 2 Q7YQW6 | Q7YQW6 bradypus tr |
| 35 | 40 | 80.0 | 445 | 2 Q7YQO7 | Q7YQO7 tapirus bai |
| 36 | 40 | 80.0 | 445 | 2 Q7YQO0 | Q7YQO0 cynocephalu |
| 37 | 40 | 80.0 | 445 | 2 Q7YR07 | Q7YR07 lemur catta |
| 38 | 40 | 80.0 | 445 | 2 Q7YR14 | Q7YR14 rhinolophus |
| 39 | 39 | 78.0 | 421 | 2 Q7TN68 | Q7TN68 glaucomys v |
| 40 | 39 | 78.0 | 432 | 2 Q7YR10 | Q7YR10 dicerops bic |
| 41 | 39 | 78.0 | 436 | 2 Q7YQW8 | Q7YQW8 nyctimene a |
| 42 | 39 | 78.0 | 438 | 2 Q7YQW7 | Q7YQW7 pteropus hy |
| 43 | 39 | 78.0 | 438 | 2 Q7YR04 | Q7YR04 roussettus a |
| 44 | 39 | 78.0 | 440 | 2 Q7YQW4 | Q7YQW4 myotis velli |
| 45 | 39 | 78.0 | 445 | 2 Q7YR08 | Q7YR08 chaetophrac |

ALIGNMENTS

| | | | | | |
|---|---|--------------|-----------|-------------------------|--|
| RESULT 1 | | | | | |
| Q7LZ77 | Q7LZ77 | PRELIMINARY; | PRT; | 275 AA. | |
| AC | Q7LZ77; | | | | |
| DT | 01-MAR-2004 (TrEMBLrel. 26, Created) | | | | |
| DT | 01-MAR-2004 (TrEMBLrel. 26, Last sequence update) | | | | |
| DT | 01-MAR-2004 (TrEMBLrel. 26, Last annotation update) | | | | |
| DB | Apolipoprotein B-100 (Fragment). | | | | |
| OS | Gallus gallus (Chicken). | | | | |
| OC | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; | | | | |
| OC | Archosauria; Aves; Neognathae; Galliformes; Phasianidae; Phasianinae; | | | | |
| OC | Gallus. | | | | |
| OX | NCBI_TaxID=9031; | | | | |
| RN | [1] | | | | |
| RP | SEQUENCE FROM N.A. | | | | |
| RX | MEDLINE=90324804; PubMed=2373961; | | | | |
| RA | Law A., Scott J.; | | | | |
| RT | "A cross-species comparison of the apolipoprotein B domain that binds | | | | |
| RL | to the LDL receptor."; | | | | |
| RL | J. Lipid Res. 31:1109-1120(1990). | | | | |
| DR | PIR; E60950; E60950. | | | | |
| FT | NON_TER | 1 | 275 | | |
| FT | NON_TER | 1 | 275 | | |
| SQ | SEQUENCE | 275 AA; | 30678 MW; | B7D8DA054E04B255 CRC64; | |
| Query Match 88.0%; Score 44; DB 2; Length 275; | | | | | |
| Best Local Similarity 90.0%; Pred. No. 1.8; Mismatches 0; Indels 0; Gaps 0; | | | | | |
| Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0; | | | | | |
| Qy | 1 TDLTRKRLK 10 | | | | |
| | | | | | |
| Db | 221 TSLTRKRLK 230 | | | | |
| RESULT 2 | | | | | |
| Q7YQN2 | Q7YQN2 | PRELIMINARY; | PRT; | 387 AA. | |
| AC | Q7YQN2; | | | | |
| DT | 01-OCT-2003 (TrEMBLrel. 25, Created) | | | | |
| DT | 01-OCT-2003 (TrEMBLrel. 25, Last sequence update) | | | | |
| DT | 01-OCT-2003 (TrEMBLrel. 25, Last annotation update) | | | | |
| DE | Apolipoprotein B 100 (Fragment). | | | | |
| GN | Name=apob-100; | | | | |
| OS | Phalanger orientalis (gray cuscus). | | | | |
| OC | Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; | | | | |
| OC | Mammalia; Metatheria; Diprotodontia; Phalangeridae; Phalanger. | | | | |
| OX | NCBI_TaxID=42473; | | | | |
| RN | [1] | | | | |
| RP | SEQUENCE FROM N.A. | | | | |
| RX | MEDLINE=22761261; PubMed=12878460; | | | | |
| RA | Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.; | | | | |
| RT | "A new phylogenetic marker, apolipoprotein B, provides compelling | | | | |
| RT | evidence for eutherian relationships."; | | | | |
| RL | Mol. Phylogenet. Evol. 28:225-240(2003). | | | | |

```
DR EMBL; AF548431; AAP97387.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 387
SQ SEQUENCE 387 AA; 43230 MW; 8300A9D7C54B42B0 CRC64;
Query Match 88.0%; Score 44; DB 2; Length 387;
Best Local Similarity 90.0%; Pred. No. 2.5;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TDLTRKRGK 10
DB 260 TSLTRKRGK 269
RESULT 3
QYQNO
ID Q7YQNO PRELIMINARY; PRT; 400 AA.
AC Q7YQNO;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apob-100;
OS Ornithorhynchus anatinus (Duckbill platypus).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Monotremata; Ornithorhynchidae; Ornithorhynchus.
OX NCBI_TaxID=9258;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548434; AAP97390.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 400
SQ SEQUENCE 400 AA; 44611 MW; DC79873CA6D01CFA CRC64;
Query Match 88.0%; Score 44; DB 2; Length 400;
Best Local Similarity 90.0%; Pred. No. 2.6;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TDLTRKRGK 10
DB 260 TSLTRKRGK 269
RESULT 4
QYQNO
ID Q7YQNO PRELIMINARY; PRT; 405 AA.
AC Q7YQNO;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apob-100;
OS Tachyglossus aculeatus (Australian echidna).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Monotremata; Tachyglossidae; Tachyglossus.
OX NCBI_TaxID=9261;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548433; AAP97389.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 1
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FT NON_TER 405
SQ SEQUENCE 405 AA; 44975 MW; 551A98557E8B081D CRC64;
Query Match 88.0%; Score 44; DB 2; Length 405;
Best Local Similarity 90.0%; Pred. No. 2.7;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TDLTRKRGK 10
DB 260 TSLTRKRGK 269
RESULT 5
QYQRS
ID Q7YQRS PRELIMINARY; PRT; 414 AA.
AC Q7YQRS;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apob-100;
OS Actus vociferans (Spix's owl monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Actinae; Actus.
OX NCBI_TaxID=57176;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548396; AAP97352.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 414
SQ SEQUENCE 414 AA; 45955 MW; EEFA8492157E1BDE CRC64;
Query Match 84.0%; Score 42; DB 2; Length 414;
Best Local Similarity 90.0%; Pred. No. 7;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 TDLTRKRGK 10
DB 258 TRLTRKRGK 267
RESULT 6
Q28473
ID Q28473 PRELIMINARY; PRT; 596 AA.
AC Q28473;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Apolipoprotein B (Fragment).
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecoidea; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RX MEDLINE=92075708; PubMed=1742325;
RA Pape M.E., Castle C.K., Murray R.W., Funk G.M., Hunt C.E.,
RA Marotti K.R., Melchior G.W.;
RT "Apo B metabolism in the cynomolgus monkey: evidence for post-
RT transcriptional regulation.";
RL Biochim. Biophys. Acta 1086:326-334(1991).
RN [2]
RP SEQUENCE FROM N.A.
RC TISSUE=Liver;
RA Murray R.;
```

RL Submitted (FEB-1992) to the EMBL/GenBank/DBJ databases.

DR EMBL; X15737; CRA33755.1; --
 DR PIR; S32802; S32802.
 KW Lipoprotein.

FT NON_TER 1 596
 FT NON_TER 596 596

SQ SEQUENCE 596 AA; 66757 MW; B13BBA74E25C3120 CRC64;
 Query Match 84.0%; Score 42; DB 2; Length 596;
 Best Local Similarity 90.0%; Pred. No. 10;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGGLK 10
 | | | | | | | | | |
 Db 226 TRLTRKRGGLK 235

RESULT 7

Q13788 PRELIMINARY; PRT; 3262 AA.
 AC Q13788;
 DT 01-NOV-1996 (TrEMBLrel. 01, Created)
 DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
 DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
 DE APOB protein (Fragment).
 GN Name=APOB;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87191999; PubMed=2883086;
 RA Carlsson P., Darnfors C., Olofsson S.O., Bjursell G.;
 RT "Analysis of the human apolipoprotein B gene; complete structure of
 the B-74 region.";
 RL Gene 49:29-51(1986).
 DR EMBL; M15421; AAA51758.1; --
 DR PIR; A27850; LPHUB.
 DR GO; GO:0005576; C:extracellular; NAS.
 DR GO; GO:0005319; F:lipid transporter activity; NAS.
 DR GO; GO:0006869; F:lipid transport; NAS.
 FT NON_TER 1
 SQ SEQUENCE 3262 AA; 370140 MW; 56603BC0618DD40D CRC64;
 Query Match 84.0%; Score 42; DB 2; Length 3262;
 Best Local Similarity 90.0%; Pred. No. 63;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGGLK 10
 | | | | | | | | | |
 Db 2084 TRLTRKRGGLK 2093

RESULT 8

APB HUMAN
 ID APB HUMAN STANDARD; PRT; 4563 AA.
 AC P04114; C00502; Q13787;
 DT 01-NOV-1986 (Rel. 03, Created)
 DT 01-NOV-1986 (Rel. 03, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Apolipoprotein B-100 precursor (Apo B-100) [Contains: Apolipoprotein
 B-48 (Apo B-48)].
 GN Name=APOB;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87016385; PubMed=3763409;
 RA Knott T.C., Wallis S.C., Powell L.M., Pease R.J., Lusis A.J.,
 RA Blackhart B., McCarthy B.J., Mahley R.W., Levy-Wilson B., Scott J.;

RT "Complete cDNA and derived protein sequence of human apolipoprotein B-
 100.";
 RL Nucleic Acids Res. 14:7501-7503(1986).
 RN [2]
 RP SEQUENCE FROM N.A., AND VARIANT GLU-4181.
 RX MEDLINE=88003974; PubMed=3652907;
 RA Ludwig E.H., Blackhart B.D., Pierotti V.R., Caiati L., Fortier C.,
 RA Knott T., Scott J., Mahley R.W., Levy-Wilson B., McCarthy B.J.;
 RT "DNA sequence of the human apolipoprotein B gene.";
 RL DNA 6:363-372(1987).
 RN [3]
 RP SEQUENCE FROM N.A., AND VARIANTS ILE-98 AND GLU-4181.
 RX MEDLINE=87008488; PubMed=3759943;
 RA Chen S.-H., Yang C.-Y., Chen P.-F., Setzer D., Tanimura M., Li W.-H.,
 RA Gotto A.M. Jr., Chan L.;
 RT "The complete cDNA and amino acid sequence of human apolipoprotein B-
 100.";
 RL J. Biol. Chem. 261:12918-12921(1986).
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87041416; PubMed=3464946;
 RA Law S.W., Grant S.M., Higuchi K., Hospattankar A.V., Lackner K.J.,
 RA Lee N., Brewer H.B. Jr.;
 RT "Human liver apolipoprotein B-100 cDNA: complete nucleic acid and
 derived amino acid sequence.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:8142-8146(1986).
 RN [5]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87161758; PubMed=3030729;
 RA Cladaras C., Hadzopoulou-Cladaras M., Nolte R.T., Atkinson D.,
 RA Zannis V.I.;
 RT "The complete sequence and structural analysis of human apolipoprotein
 B-100: relationship between apob-100 and apob-48 forms.";
 RL EMBO J. 5:3495-3507(1986).
 RN [6]
 RP SEQUENCE OF 709-906 FROM N.A.
 RX MEDLINE=85270450; PubMed=3860836;
 RA Deeb S.S., Motulsky A.G., Albers J.J.;
 RT "A partial cDNA clone for human apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 82:4983-4986(1985).
 RN [7]
 RP SEQUENCE OF 3056-3159 FROM N.A.
 RX MEDLINE=86041888; PubMed=3903660;
 RA Mehrabian M., Schumaker V.N., Fareed G.C., West R., Johnson D.F.,
 RA Kirchgessner T.G., Lin H.-C., Wang X., Ma Y., Mendiaz E., Lusis A.J.;
 RT "Human apolipoprotein B: identification of cDNA clones and
 characterization of mRNA.";
 RL Nucleic Acids Res. 13:6937-6953(1985).
 RN [8]
 RP SEQUENCE OF 1937-2018 AND 3811-4334 FROM N.A., AND VARIANT GLU-4181.
 RX MEDLINE=86093680; PubMed=3841204;
 RA Carlsson P., Olofsson S.O., Bondjers G., Darnfors C., Wiklund O.,
 RA Bjursell G.;
 RT "Molecular cloning of human apolipoprotein B cDNA.";
 RL Nucleic Acids Res. 13:8813-8826(1985).
 RN [9]
 RP SEQUENCE OF 3109-4563 FROM N.A.
 RX MEDLINE=85300528; PubMed=2994225;
 RA Knott T.J., Rall S.C. Jr., Innerarity T.L., Jacobson S.F., Urdea M.S.,
 RA Levy-Wilson B., Powell L.M., Pease R.J., Eddy R., Nakai H., Byers M.,
 RA Priestley L.M., Robertson E., Rall L.B., Betsholtz C., Shows T.B.,
 RA Mahley R.W., Scott J.;
 RT "Human apolipoprotein B: structure of carboxyl-terminal domains, sites
 of gene expression, and chromosomal localization.";
 RL Science 230:37-43(1985).
 RN [10]
 RP SEQUENCE OF 1-291 FROM N.A.
 RX MEDLINE=86149325; PubMed=3513177;
 RA Protter A.A., Hardman D.A., Schilling J.W., Miller J., Appleby V.,
 RA Chen G.C., Kirsher S.W., McEnroe G., Kane J.P.;
 RT "Isolation of a cDNA clone encoding the amino-terminal region of human
 apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:1467-1471(1986).

RN [11]
 RX SEQUENCE OF 1-1670 FROM N.A., AND VARIANT ILE-98.
 RA MEDLINE=86287319; PubMed=3461454;
 RA Protter A.A., Hardman D.A., Sato K.Y., Schilling J.W., Yamanaka M.,
 RA Hott Y.J., Hjerrild K.A., Chen G.C., Kane J.P.;
 RT "Analysis of cDNA clones encoding the entire B-26 region of human
 RT apolipoprotein B."
 RL Proc. Natl. Acad. Sci. U.S.A. 83:5678-5682(1986).
 RN [12]
 RP PARTIAL SEQUENCE, AND IDENTIFICATION OF APO-B48.
 RX MEDLINE=88018019; PubMed=3659919;
 RA Chen S.-H., Habib G., Yang C.-H., Gu Z.-W., Lee B.R., Weng S.-H.,
 RA Silberman S.R., Cai S.-J., Deslypere J.P., Rosseneu M.,
 RA Gotto A.M. Jr., Li W.-H., Chan L.;
 RT "Apolipoprotein B-48 is the product of a messenger RNA with an organ-
 RT specific in-frame stop codon."
 RL Science 238:363-366(1987).
 RN [13]
 RP DOMAINS.
 RX MEDLINE=87039351; PubMed=3773997;
 RA Knott T.C., Pease R.J., Powell L.M., Wallis S.C., Rall S.C. Jr.,
 RA Innerarity T.L., Blackhart B., Taylor W.R., Marcel Y., Milne R.,
 RA Johnson D., Fuller M., Lusis A.J., McCarthy B.J., Mahley R.W.,
 RA Levy-Wilson B., Scott J.;
 RT "Complete protein sequence and identification of structural domains of
 RT human apolipoprotein B."
 RL Nature 323:734-738(1986).
 RN [14]
 RP DOMAINS.
 RA Yang C.-Y., Chen S.-H., Gianturco S.H., Bradley W.A., Sparrow J.T.,
 RA Tanimura M., Li W.-H., Sparrow D.A., Deloof H., Rosseneu M.,
 RA Lee F.-S., Gu Z.-W., Gotto A.M. Jr., Chan L.;
 RT "Sequence, structure, receptor-binding domains and internal repeats of
 RT human apolipoprotein B-100."
 RL Nature 323:738-742(1986).
 RN [15]
 RP CALCIUM-BINDING DATA.
 RX MEDLINE=86242245; PubMed=3087360;
 RA Dashti N., Lee D.M., Mok T.;
 RT "Apolipoprotein B is a calcium binding protein."
 RL Biochem. Biophys. Res. Commun. 137:493-499(1986).
 RN [16]
 RP PALMITOYLATION OF CYS-1112.
 RX MEDLINE=20143590; PubMed=10679026;
 RA Zhao Y., McCabe J.B., Vance J., Berthiaume L.G.;
 RT "Palmitoylation of apolipoprotein B is required for proper
 RT intracellular sorting and transport of cholesterol esters and
 RT triglycerides."
 RL Mol. Biol. Cell 11:721-734(2000).
 RN [17]
 RP VARIANT SER-4338.
 RX MEDLINE=91071750; PubMed=1979313;
 RA Navajas M., Laurent A.-M., Moreel J.-F., Ragab A., Cambou J.-P.,
 RA Cuny G., Cambien F., Roizes G.;
 RT "Detection by denaturing gradient gel electrophoresis of a new
 RT polymorphism in the apolipoprotein B gene."
 RL Hum. Genet. 86:91-93(1990).
 RN [18]
 RP VARIANT FDB GLN-3527.
 RX MEDLINE=89098975; PubMed=2563166;
 RA Sorla L.F., Ludwig E.H., Clarke H.R.G., Vega G.L., Grundy S.M.,
 RA McCarthy B.J.;
 RT "Association between a specific apolipoprotein B mutation and familial
 RT defective apolipoprotein B-100."
 RL Proc. Natl. Acad. Sci. U.S.A. 86:587-591(1989).
 RN [19]
 RP VARIANT LEU-2739.
 RX MEDLINE=91016974; PubMed=2216805;
 RA Huang L.-S., Gavish D., Breslow J.L.;
 RT "Sequence polymorphism in the human apoB gene at position 8344."
 RL Nucleic Acids Res. 18:5922-5922(1990).
 RN [20]
 RP VARIANT FDB CYS-3558.

RX MEDLINE=95190020; PubMed=7883971;
 RA Pullinger C.R., Hennessey L.K., Chatterton J.E., Liu W., Love J.A.,
 RA Mendel C.M., Frost P.H., Malloy M.J., Schumaker V.N., Kane J.P.;
 RT "Familial ligand-defective apolipoprotein B. Identification of a new
 RT mutation that decreases LDL receptor binding affinity."
 RL J. Clin. Invest. 95:1225-1234(1995).
 RN [21]
 RP VARIANTS LEU-1437; SER-1914; LYS-2566; THR-3121; ALA-3945; MET-4128
 RP AND THR-4481.
 RX MEDLINE=97044521; PubMed=8889592;
 RA Poiret O., Ricard S., Behague I., Souriau C., Evans A.E.,
 RA Arveiler D., Marques-Vidal P., Luc G., Roizes G., Cambien F.;
 RT "Detection of new variants in the apolipoprotein B (Apo B) gene by
 RT PCR-SSCP."
 RL Hum. Mutat. 8:282-285(1996).
 RN [22]
 RP VARIANTS FDB GLN-3527 AND CYS-3558.
 RX MEDLINE=97403938; PubMed=9259199;
 RA Rabes J.P., Varret M., Saint-Jore B., Erlich D., Jondeau G.,
 RA Krempf M., Giraudet P., Junien C., Boileau C.;
 RT "Familial ligand-defective apolipoprotein B-100: simultaneous
 RT detection of the ARG3500-->GLN and ARG3531-->CYS mutations in a French
 RT population."
 RL Hum. Mutat. 10:160-163(1997).
 RN [23]
 RP VARIANTS SER-1914; ARG-1923; LEU-2739; ASP-3319; THR-3427; GLN-3432
 RP AND ILE-3921.
 RX MEDLINE=98141125; PubMed=9490296;
 RA Leren T.P., Bakken K.S., Hoel V., Hjermann I., Berg K.;
 RT "Screening for mutations of the apolipoprotein B gene causing
 RT hypocholesterolemia."
 RL Hum. Genet. 102:44-49(1998).
 CC -1- FUNCTION: Apolipoprotein B is a major protein constituent of
 CC chylomicrons (apo B-48), LDL (apo B-100) and VLDL (apo B-100). Apo
 CC B-100 functions as a recognition signal for the cellular binding
 CC and internalization of LDL particles by the apoB/E receptor.
 CC -1- SUBCELLULAR LOCATION: Secreted.
 Query Match 84.0%; Score 42; DB 1; Length 4563;
 Best Local Similarity 90.0%; Pred. No. 91;
 Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TDLTKRGLK 10
 | | | | | | | | | |
 Db 3385 TELTKRGLK 3394
 RESULT 9
 Q7Z600
 ID Q7Z600 PRELIMINARY; PRT; 4563 AA.
 AC Q7Z600;
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Apolipoprotein B (including Ag(X) antigen).
 GN Name=APOB;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
 RA Ahearn M.O., Kuldane S.A., Rajkumar N., Toth E.J., Yi Q.,
 RA Nickerson D.A.;
 RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AY324608; AAF72970.1; -
 DR GO; GO:0005319; F:lipid transporter activity; IEA.
 DR GO; GO:0006869; P:lipid transport; IEA.
 DR InterPro; IPR009454; DUF1081.
 DR InterPro; IPR001747; Lipid_transprt_N.
 DR Pfam; PF06448; DUF1081; 1.
 DR Pfam; PF01347; Vitellogenin_N; 1.

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DR SMART; SM00638; LPD_N; 1.
KW Lipoprotein.
SQ SEQUENCE 4563 AA; 51553 MW; 030B34167CEDC63C CRC64;

Query Match      84.0%; Score 42; DB 2; Length 4563;
Best Local Similarity 90.0%; Pred. No. 91;
Matches 9; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGK 10
Db 3385 TRLTRKRGK 3394

RESULT 10
Q7YQ00 PRELIMINARY; PRT; 263 AA.
AC Q7YQ00;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Procavia capensis (Cape hyrax) (Rock dassie).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Hyracoidea; Procaviidae; Procavia.
OX NCBI_TaxID=9813;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548411; AAP97367.1; -.
KW Lipoprotein.
FT NON_TER 1 1
FT NON_TER 263 263
SQ SEQUENCE 263 AA; 29532 MW; 536CF6149C1D062A CRC64;

Query Match      80.0%; Score 40; DB 2; Length 263;
Best Local Similarity 80.0%; Pred. No. 11;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGK 10
Db 206 SSLTRKRGK 215

RESULT 11
Q7M2U9 PRELIMINARY; PRT; 274 AA.
AC Q7M2U9;
DT 01-MAR-2004 (TrEMBLrel. 26, Created)
DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Apolipoprotein B-100 (Fragment).
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90324804; PubMed=2373961;
RA Law A., Scott J.;
RT "A cross-species comparison of the apolipoprotein B domain that binds
RT to the LDL receptor.";
RL J. Lipid Res. 31:1109-1120(1990).
DR PIR; A60950; A60950.
FT NON_TER 1 1
FT NON_TER 274 274
SQ SEQUENCE 274 AA; 30505 MW; CA1E1BE360AAB8F2 CRC64;

Query Match      80.0%; Score 40; DB 2; Length 274;
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Best Local Similarity 80.0%; Pred. No. 12;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGK 10
Db 221 SSLTRKRGK 230

RESULT 12
Q7YQP9 PRELIMINARY; PRT; 304 AA.
AC Q7YQP9;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Echinops telfairi (Lesser hedgehog tenrec).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Insectivora; Tenrecidae; Tenrecinae; Echinops.
OX NCBI_TaxID=9371;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548412; AAP97368.1; -.
KW Lipoprotein.
FT NON_TER 1 1
FT NON_TER 304 304
SQ SEQUENCE 304 AA; 34264 MW; 468F4409260D6358 CRC64;

Query Match      80.0%; Score 40; DB 2; Length 304;
Best Local Similarity 80.0%; Pred. No. 13;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TDLTRKRGK 10
Db 176 SSLTRKRGK 185

RESULT 13
Q7YQN8 PRELIMINARY; PRT; 314 AA.
AC Q7YQN8;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Ictonyx striatus (striped polecat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Carnivora; Fissipedia; Mustelidae; Mustelinae;
OC Ictonyx.
OX NCBI_TaxID=55050;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548425; AAP97381.1; -.
KW Lipoprotein.
FT NON_TER 1 1
FT NON_TER 314 314
SQ SEQUENCE 314 AA; 34719 MW; 3E5B34E780F0039E CRC64;

Query Match      80.0%; Score 40; DB 2; Length 314;
Best Local Similarity 80.0%; Pred. No. 14;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
```

QY 1 TDLTRKRGK 10
: |||||
Db 162 SSLTRKRGK 171

Db : |||||
162 SSLTRKRGK 171

Search completed: January 13, 2005, 01:51:03
Job time : 79.0328 secs

RESULT 14

Q7YQP3
ID Q7YQP3 PRELIMINARY; PRT; 316 AA.
AC Q7YQP3;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Nandinia binotata (African palm civet).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Carnivora; Fissipedia; Viverridae; Nandiniinae;
OC Nandinia.
OX NCBI_TaxID=71115;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548420; AAP97376.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 316
SQ SEQUENCE 316 AA; 34540 MW; C04896B0E17562AE CRC64;

Query Match 80.0%; Score 40; DB 2; Length 316;
Best Local Similarity 80.0%; Pred. No. 14;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRGK 10
: |||||
Db 160 SSLTRKRGK 169

RESULT 15

Q7YQN9
ID Q7YQN9 PRELIMINARY; PRT; 318 AA.
AC Q7YQN9;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apoB-100;
OS Zalophus californianus (California sealion).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Carnivora; Pinnipedia; Otariidae; Zalophus.
OX NCBI_TaxID=9704;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548424; AAP97380.1; -.
KW Lipoprotein.
FT NON_TER 1
FT NON_TER 318
SQ SEQUENCE 318 AA; 34888 MW; C04E7ECBA8E64C96 CRC64;

Query Match 80.0%; Score 40; DB 2; Length 318;
Best Local Similarity 80.0%; Pred. No. 14;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TDLTRKRGK 10

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 69.4918 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-15

Perfect score: 44

Sequence: 1 TRLTRKGLK 9

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_23Sep04:*

1: Geneseq1980s:*

2: Geneseq1990s:*

3: Geneseq2000s:*

4: Geneseq2001s:*

5: Geneseq2002s:*

6: Geneseq2003as:*

7: Geneseq2003bs:*

8: Geneseq2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|-------|--------------------|
| 1 | 44 | 100.0 | 9 | 2 | AAy30696 Apo-B100 |
| 2 | 41 | 93.2 | 9 | 2 | AAy30694 Apo-B100 |
| 3 | 38 | 86.4 | 9 | 2 | AAy30695 Apo-B100 |
| 4 | 34 | 77.3 | 601 | 6 | ADA47994 Rice prot |
| 5 | 34 | 77.3 | 798 | 8 | ABU02213 S. pneumo |
| 6 | 34 | 77.3 | 798 | 8 | ADK46589 Streptoco |
| 7 | 34 | 77.3 | 804 | 3 | AAy55803 S. pneumo |
| 8 | 33.5 | 76.1 | 10 | 2 | AAy30690 Apo-B100 |
| 9 | 33.5 | 76.1 | 10 | 2 | AAy30691 Apo-B100 |
| 10 | 33.5 | 76.1 | 11 | 2 | AAW57205 Apo B bin |
| 11 | 33.5 | 76.1 | 13 | 2 | AAW57207 Apo B 100 |
| 12 | 33.5 | 76.1 | 15 | 2 | AAW41261 Apolipop |
| 13 | 33.5 | 76.1 | 15 | 2 | AAW96892 ApoB-100 |
| 14 | 33.5 | 76.1 | 20 | 6 | ABJ37575 Heparin b |
| 15 | 33.5 | 76.1 | 22 | 2 | AAW57208 Apo B 100 |
| 16 | 33.5 | 76.1 | 22 | 2 | AAW57209 Apo B 100 |
| 17 | 33.5 | 76.1 | 34 | 5 | AAE14541 Human apo |
| 18 | 33.5 | 76.1 | 36 | 2 | AAW96876 Nucleic a |
| 19 | 33.5 | 76.1 | 37 | 2 | AAW64587 Human apo |
| 20 | 33.5 | 76.1 | 51 | 2 | AAW96845 Nucleic a |
| 21 | 33.5 | 76.1 | 343 | 4 | ABB37687 Peptide # |
| 22 | 33.5 | 76.1 | 343 | 4 | ABG52504 Human liv |
| 23 | 33.5 | 76.1 | 377 | 2 | AAr72704 Human apo |
| 24 | 33.5 | 76.1 | 377 | 2 | AAr34031 Sequence |
| 25 | 33.5 | 76.1 | 2463 | 8 | ADJ57400 Human apo |

| | | | | | |
|----|------|------|------|---|--------------------|
| 26 | 33.5 | 76.1 | 3923 | 2 | AAy31237 Human Apo |
| 27 | 33.5 | 76.1 | 4536 | 2 | AAW41262 Apolipop |
| 28 | 33.5 | 76.1 | 4536 | 2 | AAW96826 Amino aci |
| 29 | 33.5 | 76.1 | 4560 | 5 | AAU98981 Human apo |
| 30 | 33.5 | 76.1 | 4561 | 7 | ADD48677 Human Pro |
| 31 | 33.5 | 76.1 | 4563 | 5 | AAO15893 Human apo |
| 32 | 33.5 | 76.1 | 4563 | 6 | ABR40253 Human ali |
| 33 | 33.5 | 76.1 | 4563 | 6 | ABU79140 Apolipop |
| 34 | 33.5 | 76.1 | 4563 | 7 | ADf43408 Apolipop |
| 35 | 33.5 | 76.1 | 4563 | 8 | ADH18871 Human apo |
| 36 | 33.5 | 76.1 | 4563 | 8 | ADH18870 Human apo |
| 37 | 33.5 | 76.1 | 4563 | 8 | ADO33445 Human apo |
| 38 | 33.5 | 76.1 | 4563 | 8 | ADO33447 Human apo |
| 39 | 33.5 | 76.1 | 4590 | 4 | AAU33184 Novel hum |
| 40 | 33 | 75.0 | 11 | 2 | AAy30700 Apo-B100 |
| 41 | 33 | 75.0 | 11 | 2 | AAy30698 Apo-B100 |
| 42 | 33 | 75.0 | 147 | 5 | ABB49220 Listeria |
| 43 | 33 | 75.0 | 150 | 7 | ADC89104 Ribosomal |
| 44 | 33 | 75.0 | 151 | 7 | ADC88027 Ribosomal |
| 45 | 33 | 75.0 | 154 | 7 | ADM26985 Hyperther |

ALIGNMENTS

RESULT 1

AAy30696

ID AAY30696 standard; peptide; 9 AA.

XX AC AAY30696;

XX 17-NOV-1999 (first entry)

XX DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

XX KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;

XX KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

XX OS Synthetic.

OS Homo sapiens.

XX WO9946598-A1.

XX 16-SEP-1999.

XX 05-MAR-1999; 99WO-US004805.

XX 10-MAR-1998; 98US-0077618P.

XX (REGC) UNIV CALIFORNIA.

XX Innerarity TL, Boren JOS;

XX WPI; 1999-551509/46.

PT Identifying compounds which affect binding of low density lipoprotein with proteoglycan, used for, e.g. obtaining compounds for reducing atherosclerosis.

PT Claim 17; Page 57; 70pp; English.

XX AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan receptor mutations. They were created to identify compounds which modulate atherosclerosis. The peptides are derived from amino acids 3358 to 3367 of apoB100. The method comprises detecting compounds which affect low density lipoprotein (LDL) binding with proteoglycan (PG). The method can be used for identifying compounds which disrupt LDL-PG binding without inhibiting LDL receptor binding. Such compounds can be used to reduce or prevent the formation of atherosclerotic lesions and prevent atherosclerosis. The transgenic non-human animals and mammals which express human apo-B100 can be used as an in vivo model system for the study of atherosclerosis, and in vivo assay methods for identifying compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 9 AA;

Query Match 100.0%; Score 44; DB 2; Length 9;
 Best Local Similarity 100.0%; Pred. No. 1.7e+06;
 Matches 9; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKGLK 9
 DB 1 TRLTRKGLK 9

RESULT 2

AAAY30694
 ID AAY30694 standard; peptide; 9 AA.

AC AAY30694;

DT 17-NOV-1999 (first entry)

DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

OS Synthetic.

OS Homo sapiens.

PN WO9946598-A1.

PD 16-SEP-1999.

PF 05-MAR-1999; 99WO-US004805.

PR 10-MAR-1998; 98US-0077618P.

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PI Innerarity TL, Boren JOS;

PS WPI; 1999-551509/46.

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PS Claim 17; Page 57; 70pp; English.

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 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 9 AA;

Query Match 93.2%; Score 41; DB 2; Length 9;
 Best Local Similarity 88.9%; Pred. No. 1.7e+06;
 Matches 8; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKGLK 9
 DB 1 TRLTRKGLK 9

RESULT 3

AAAY30695
 ID AAY30695 standard; peptide; 9 AA.

XX AAY30695;

DT 17-NOV-1999 (first entry)

DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

OS Synthetic.

OS Homo sapiens.

PN WO9946598-A1.

PD 16-SEP-1999.

PF 05-MAR-1999; 99WO-US004805.

PR 10-MAR-1998; 98US-0077618P.

PA (REGC) UNIV CALIFORNIA.

PI Innerarity TL, Boren JOS;

DR WPI; 1999-551509/46.

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 with proteoglycan, used for, e.g. obtaining compounds for reducing
 atherosclerosis.

PS Claim 17; Page 57; 70pp; English.

CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

SQ Sequence 9 AA;

Query Match 86.4%; Score 38; DB 2; Length 9;
 Best Local Similarity 77.8%; Pred. No. 1.7e+06;
 Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKGLK 9

```

Db      |||:||||
      1  TRLTRKGLK 9

RESULT 4
ADA47994
XX  ID  ADA47994 standard; protein; 601 AA.
XX  AC
XX  AD  ADA47994;
XX  DT  20-NOV-2003 (first entry)
XX  DE  Rice protein conferring disease resistance in plants.
XX  KW  disease resistance; pathogen tolerance; plant pathogen; plant; rice.
XX  KW  Oryza sativa.
XX  OS
XX  FN  WO2003000906-A2.
XX  PD  03-JAN-2003.
XX  PF  21-JUN-2002; 2002WO-IB002453.
XX  PR  22-JUN-2001; 2001US-0300112P.
XX  PR  26-SEP-2001; 2001US-0352277P.
XX  PR  22-MAR-2002; 2002US-0366535P.
XX  SYGN ) SYNGENTA PARTICIPATIONS AG.
XX  PI  Glazebrook J, Briggs S, Cooper B, Goff SA, Moughamer T;
XX  PI  Katagiri F, Kreps J, Provart N, Ricke D, Zhu T;
XX  DR  WPI; 2003-184052/18.
XX  DR  N-PSDB; ADA47993.
XX  PT  New polynucleotide comprising a plant nucleotide sequence having an open
XX  PT  reading frame that encodes a polypeptide associated with disease
XX  PT  resistance, useful for conferring resistance or tolerance to a plant
XX  PT  pathogen.
XX  PS  Claim 10; SEQ ID NO 64; 299pp; English.
XX  CC  The invention relates to a novel isolated polynucleotide comprising a
XX  CC  plant nucleotide sequence having an open reading frame that encodes a
XX  CC  polypeptide associated with disease resistance or its fragment having
XX  CC  substantially the same activity as the full-length polypeptide. The
XX  CC  polynucleotide of the invention is useful for conferring resistance or
XX  CC  tolerance to a plant pathogen. The present sequence represents a protein
XX  CC  conferring disease resistance used in the invention.
XX  SQ  Sequence 601 AA;

Query Match      77.3%; Score 34; DB 6; Length 601;
Best Local Similarity 87.5%; Pred. No. 2.2e+02;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy      1  TRLTRKGL 8
      ||| ||| |||
Db      263  TRLTRSGL 270

RESULT 5
ABU02213
XX  ID  ABU02213 standard; protein; 798 AA.
XX  AC
XX  AD  ABU02213;
XX  DT  23-OCT-2003 (revised)
XX  DT  11-FEB-2003 (first entry)
XX  DE  S. pneumoniae type 4 strain protein from coding region #1791.
XX  KW  Bacterial meningitis; pneumonia; sepsis; otitis media; ear infection;
XX  KW  antiinflammatory; antibacterial; immunostimulant; auditory; respiratory;
XX  KW  gene therapy; vaccine.
XX  OS  Streptococcus pneumoniae; type 4 strain.
XX  FN  WO200277021-A2.
XX  PD  03-OCT-2002.
XX  PF  27-MAR-2002; 2002WO-IB002163.
XX  PR  27-MAR-2001; 2001GB-00007658.
XX  PA  (CHIR-) CHIRON SPA.
XX  PA  (GENO-) INST GENOMIC RES.
XX  PI  Masignani V, Tettelin H, Fraser C;
XX  DR  WPI; 2003-040579/03.
XX  DR  N-PSDB; ABX07503.
XX  PT  New proteins and nucleic acid molecules from Streptococcus pneumoniae,
XX  PT  useful as medicaments for treating or preventing a disease or infection
XX  PT  due to streptococcus bacteria, such as pneumonia, sepsis, otitis media or
XX  PT  ear infection.
XX  PS  Claim 1; SEQ ID NO 3582; 56pp; English.
XX  CC  The invention relates to a protein comprising or having at least 50%
XX  CC  identity to any of the 2469 amino acid sequences, identified in the
XX  CC  specification (available on a computer readable format), or its fragment,
XX  CC  expressed from 2469 of 2489 identified DNA coding regions from the
XX  CC  Streptococcus pneumoniae type 4 strain genomic sequence appearing as
XX  CC  ABS56454. Also included are an antibody which binds one of the proteins,
XX  CC  treating a patient by administering the protein, DNA or antibody (in a
XX  CC  composition), a kit comprising first and second primers, which are the
XX  CC  nucleic acid cited above or fragments between nucleotides 8-100 of a
XX  CC  sequence not defined in the specification, for amplifying a target
XX  CC  sequence contained within a Streptococcus nucleic acid sequence, where
XX  CC  the first primer is substantially complementary to the target sequence and
XX  CC  the second primer is substantially complementary to the complement of
XX  CC  the target sequence, and where the parts of the primers having
XX  CC  substantial complementarity define the termini of the target sequence to
XX  CC  be amplified, assay comprising contacting a test compound with the
XX  CC  protein, and determining whether the test compound binds to the protein
XX  CC  encoding the proteins has been rendered inactive. The proteins, nucleic
XX  CC  acid molecules, antibody and compositions are useful as medicaments for
XX  CC  treating or preventing a disease or infection due to streptococcus
XX  CC  bacteria, particularly S. pneumoniae, such as pneumonia, sepsis, otitis
XX  CC  media or ear infection. They are also useful in developing vaccines,
XX  CC  diagnostics and antibiotics. The methods are useful for identifying
XX  CC  immunodominant proteins. The present sequence is one of the 2469 proteins
XX  CC  expressed by the identified coding regions from the genomic sequence.
XX  CC  Note: The sequence data for this patent did not form part of the printed
XX  CC  specification, but was obtained in electronic format directly from WIPO
XX  CC  at ftp.wipo.int/pub/published_pct_sequences. (Updated on 23-OCT-2003 to
XX  CC  standardise OS field)
XX  SQ  Sequence 798 AA;

Query Match      77.3%; Score 34; DB 6; Length 798;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy      2  RLTRKGL 8
      ||| ||| |||
Db      156  RLTRKGL 162

RESULT 6
ADK46589

```

```

ID ADK46589 standard; protein; 798 AA.
XX
AC ADK46589;
XX
DT 20-MAY-2004 (first entry)
XX
DE Streptococcus pneumoniae protein; Seq ID No 3104.
XX
KW Antibacterial; Gene therapy; Vaccine; Streptococcus pneumoniae.
XX
OS Streptococcus pneumoniae.
XX
PN US6699703-B1.
XX
PD 02-MAR-2004.
XX
PF 26-MAY-2000; 2000US-00583110.
XX
PR 02-JUL-1997; 97US-0051553P.
PR 12-MAY-1998; 98US-0085131P.
PR 30-JUN-1998; 98US-00107433.
XX
PA (GENO-) GENOME THERAPEUTICS CORP.
XX
PI Doucette-Stamm L, Bush D, Zeng Q, Opperman T, Houseweart CE;
XX
DR WPI; 2004-212399/20.
XX
DR N-PSDB; ADK43928.
XX
PT New nucleic acid molecules and polypeptides useful for diagnosing,
PT preventing and treating pathological conditions resulting from bacterial
PT infection, e.g. Streptococcus pneumoniae infection, and in drug
PT screening.
XX
PS Disclosure; SEQ ID NO 3104; 301pp; English.
XX
CC The invention relates to isolated Streptococcus pneumoniae nucleic acids
CC and polypeptides. The nucleic acids and proteins are useful for
CC diagnosing, preventing and treating pathological conditions resulting
CC from bacterial infection, such as S. pneumoniae infection. These may also
CC be used for drug screening procedures. The present sequence represents a
CC Streptococcus pneumoniae polypeptide of the invention. Note: The sequence
CC data for this patent did not appear in the printed specification but was
CC obtained in electronic format directly from USPTO at
CC seqdata.uspto.gov/sequence.html.
XX
SQ Sequence 798 AA;

Query Match 77.3%; Score 34; DB 8; Length 798;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 RLTRKGL 8
Db 156 RLTRKGL 162

RESULT 7
AAV55803
ID AAV55803 standard; protein; 804 AA.
XX
AC AAV55803;
XX
DT 28-FEB-2000 (first entry)
XX
DE S. pneumoniae priA polypeptide.
XX
KW priA polypeptide; microbial disease; vaccine; microbial infection;
KW Streptococcus pneumoniae; antibacterial.
XX
OS Streptococcus pneumoniae.
XX
PN WO9961453-A2.

XX 02-DEC-1999.
XX
PD 22-APR-1999; 99WO-US008771.
XX
PR 27-APR-1998; 98US-00067091.
XX
PA (SMIK ) SMITHKLINE BEECHAM CORP.
XX
PI Mcdevitt D, Shilling L, Warren RL, St John A;
XX
DR WPI; 2000-062670/05.
XX
DR N-PSDB; AA239569, AA239572.
XX
PT New isolated priA polypeptides, useful for screening antibacterial
PT compounds.
XX
PS Claim 1; Page 4-5; 68pp; English.
XX
CC This represents a S. pneumoniae priA polypeptide. The priA polypeptides
CC and polynucleotides are useful for the treatment of microbial diseases
CC (especially in the form of vaccines) and the methods are useful for
CC identifying agonists and antagonists. The polypeptides are also useful
CC for relating to diagnostic assays for detecting diseases associated with
CC microbial infections (especially infections by Streptococcus pneumoniae)
CC and conditions associated with such infections and assays for detecting
CC priA expression or activity. The polypeptides are useful in the discovery
CC and development of antibacterial compounds. The encoded protein upon
CC expression can be used as a target for screening of antibacterial drugs
XX
SQ Sequence 804 AA;

Query Match 77.3%; Score 34; DB 3; Length 804;
Best Local Similarity 100.0%; Pred. No. 3e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 RLTRKGL 8
Db 162 RLTRKGL 168

RESULT 8
AAV30690
ID AAV30690 standard; peptide; 10 AA.
XX
AC AAV30690;
XX
DT 17-NOV-1999 (first entry)
XX
DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS Synthetic.
XX
OS Homo sapiens.
XX
PN WO9946598-A1.
XX
PD 16-SEP-1999.
XX
PF 05-MAR-1999; 99WO-US004805.
XX
PR 10-MAR-1998; 98US-0077618P.
XX
PA (REGC ) UNIV CALIFORNIA.
XX
PI Innerarity TL, Boren JOS;
XX
DR WPI; 1999-551509/46.
XX
PT Identifying compounds which affect binding of low density lipoprotein
PT with proteoglycan, used for, e.g. obtaining compounds for reducing

```

PT atherosclerosis.
 PS Claim 17; Page 57; 70pp; English.
 XX
 CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX
 SQ Sequence 10 AA;
 Query Match 76.1%; Score 33.5; DB 2; Length 10;
 Best Local Similarity 90.0%; Pred. No. 3.8;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRK-CLK 9
 DB 1 TRLTRKEGLK 10
 RESULT 9
 AAY30691
 ID AAY30691 standard; peptide; 10 AA.
 AC AAY30691;
 XX
 DT 17-NOV-1999 (first entry)
 XX
 DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
 XX
 KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 FN WO9946598-A1.
 XX
 PD 16-SEP-1999.
 XX
 PF 05-MAR-1999; 98WO-US004805.
 XX
 PR 10-MAR-1998; 98US-0077618P.
 XX
 PA (REGC) UNIV CALIFORNIA.
 XX
 FI Innerarity TL, Boren JOS;
 XX
 DR WPI; 1999-551509/46.
 XX
 XX Identifying compounds which affect binding of low density lipoprotein
 PT with proteoglycan, used for, e.g. obtaining compounds for reducing
 PT atherosclerosis.
 XX
 PS Claim 17; Page 57; 70pp; English.
 XX
 CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
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 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX
 SQ Sequence 10 AA;
 Query Match 76.1%; Score 33.5; DB 2; Length 10;
 Best Local Similarity 90.0%; Pred. No. 3.8;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRK-CLK 9
 DB 1 TRLTRKDGK 10
 RESULT 10
 AAW57205
 ID AAW57205 standard; peptide; 11 AA.
 AC AAW57205;
 XX
 DT 03-AUG-1998 (first entry)
 XX
 DE Apo B binding site peptide 2.
 XX
 KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 OS Synthetic.
 XX
 FN WO9813385-A2.
 XX
 PD 02-APR-1998.
 XX
 PF 25-SEP-1997; 97WO-GB002610.
 XX
 PR 27-SEP-1996; 96GB-00020153.
 XX
 PA (UYST) UNIV STRATHCLYDE.
 XX
 FI Halbert GW, Owens MD, Baillie G;
 XX
 DR WPI; 1998-230637/20.
 XX
 PT Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 PS Claim 12; Page 52; 73pp; English.
 XX
 CC The present sequence represents a specifically claimed Apo B binding site
 CC peptide which can be used as a component of a non-naturally occurring,
 CC receptor-competent low density lipoprotein (LDL) particle of the present
 CC invention. The LDL particle comprises at least 1 peptide component that
 CC has at least 1 binding site for an apo B protein receptor and at least 1
 CC lipophilic substituent. Also described in the invention are peptides
 CC containing an apo B binding sequence with at least 70% identity with
 CC sequences: KAEYKKNKRRH (1) or TRLTRKRGK (2), or their dimers. Non-
 CC naturally occurring, receptor-competent LDL particles are useful as: (i)
 CC drug-targeting vectors for delivering anticancer drugs to cancer cells

CC that express an apo B protein receptor, and (ii) additives for cell
 CC culture media especially as growth supplements. Non-naturally occurring,
 CC receptor-competent LDL particles do not require the complete apo B
 CC sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor

XX
 SQ Sequence 11 AA;
 Query Match 76.1%; Score 33.5; DB 2; Length 11;
 Best Local Similarity 90.0%; Pred. No. 4.2;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Oy 1 TRLTRK-GLK 9
 |||||
 Db 2 TRLTRKGLK 11

RESULT 11
 AAW57207
 ID AAW57207 standard; peptide; 13 AA.
 AC AAW57207;
 XX
 XX 03-AUG-1998 (first entry)
 DT Apo B 100 binding site peptide analogue peptide B.
 DE
 XX
 XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 XX Synthetic.
 OS
 XX
 XX Key Location/Qualifiers
 FH Modified-site 1 /note= "attached to retinoic acid"
 FT
 FT
 XX WO9813385-A2.
 XX
 XX 02-APR-1998.
 PD
 XX
 XX 25-SEP-1997; 97WO-GB002610.
 XX
 XX 27-SEP-1996; 96GB-00020153.
 PR
 XX (UVST) UNIV STRATHCLYDE.
 PA
 XX Halbert GW, Owens MD, Baillie G;
 PI WPI; 1998-230637/20.
 DR
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 XX Claim 13; Fig 7; 73pp; English.
 PS
 XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or ITRTRKGLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor

SQ Sequence 13 AA;
 Query Match 76.1%; Score 33.5; DB 2; Length 13;
 Best Local Similarity 90.0%; Pred. No. 5;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Oy 1 TRLTRK-GLK 9
 |||||
 Db 3 TRLTRKGLK 12

RESULT 12
 AAW41261
 ID AAW41261 standard; peptide; 15 AA.
 XX
 AC AAW41261;
 XX
 XX 19-MAY-1998 (first entry)
 DT Apolipoprotein B-100 fragment.
 DE
 XX
 XX Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
 KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
 KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
 KW prothrombinase complex.
 XX
 XX Synthetic.
 OS Homo sapiens.
 XX
 XX WO9743311-A1.
 PN
 XX 20-NOV-1997.
 PD
 XX
 XX 09-MAY-1997; 97WO-GB001255.
 PF
 XX 09-MAY-1996; 96GB-00009702.
 PR
 XX (UNLO) ROYAL FREE HOSPITAL SCHOOL MED.
 PA
 XX Bruckdorfer KR, Ettelaie C;
 PI WPI; 1998-008798/01.
 DR
 XX
 XX Peptide fragments of apo:lipo:protein B-100 with anticoagulant activity -
 PT used for treating or preventing coagulation, inhibiting angiogenesis,
 PT cell differentiation and apoptosis.
 XX
 XX Disclosure; Page 22; 60pp; English.
 PS
 XX This sequence is an example of the peptide of the invention. It has the
 CC formula (I), or their variants with one or more internal deletions,
 CC insertions or substitutions, while retaining anti-coagulant properties of
 CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KNKRHS-X2-T-22 (I) X1 = S or
 CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
 CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
 CC aa. Compositions containing the peptide are used for simultaneous,
 CC separate or sequential treatment of cancer, particularly to prevent
 CC metastatic spread. They are also used to inhibit thromboplastin-mediated
 CC processes, specifically to prevent or reduce blood coagulation (e.g.
 CC during or after surgery or in cases of heart attack, stroke etc.) and to
 CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
 CC which is active as such or as part of a 98-aa peptide, inhibits
 CC activation of the prothrombinase complex; and prevents activation of
 CC factor VII on the surface of thromboplastin and of platelets by thrombin.
 CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
 CC smaller than apoB-100, they act more quickly

SQ Sequence 15 AA;
 Query Match 76.1%; Score 33.5; DB 2; Length 15;
 Best Local Similarity 90.0%; Pred. No. 5.8;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRK-GLK 9
 Db ||||| |||
 1 TRLTRKRLGLK 10

RESULT 13
 AAW96892
 ID AAW96892 standard; peptide; 15 AA.
 XX AC
 XX AC AAW96892;
 XX DT 22-APR-1999 (first entry)
 XX DE ApoB-100 nuclear localisation signal sequence, residues 3353-3367.
 XX KW Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
 XX OS Homo sapiens.
 XX PN WO9856938-A1.
 XX PD 17-DEC-1998.
 XX PF 10-JUN-1998; 98WO-US011927.
 XX PR 13-JUN-1997; 97US-00874807.
 XX PR 14-MAY-1998; 98US-00079030.
 XX PA (BAYU) BAYLOR COLLEGE MEDICINE.
 XX PI Guevata JG, Hoogveen RC, Moore JP;
 XX DR WPT; 1999-070331/06.
 XX PT Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 XX used for delivering nucleic acid to cells for gene therapy and antisense
 XX treatment.
 XX PS Claim 19; Fig 13D; 293pp; English.
 XX CC AAW96878-97 represent nuclear localisation signal sequence derived from
 CC human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein
 CC component of very-low density lipoproteins (VLDL), intermediate density
 CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The
 CC present sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis
 XX SQ Sequence 15 AA;

Query Match 76.1%; Score 33.5; DB 2; Length 15;
 Best Local Similarity 90.0%; Pred. No. 5.8;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRK-GLK 9
 Db ||||| |||
 6 TRLTRKRLGLK 15

RESULT 14
 ABJ37575
 ID ABJ37575 standard; peptide; 20 AA.
 XX AC ABJ37575;
 XX XX

DT 10-MAY-2003 (first entry)
 XX Heparin binding peptide sequence #28.
 DE KW Cystostatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;
 XX KW cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;
 KW rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.
 XX OS Unidentified.
 XX PN WO2003007689-A2.
 XX PD 30-JAN-2003.
 XX PF 22-JUL-2002; 2002WO-US023419.
 XX PR 20-JUL-2001; 2001US-0306726P.
 XX PA (ETHZ-) ETH ZUERICH.
 XX PA (UYZU-) UNIV ZURICH.
 XX PI Hubbell JA, Schoenmakers R, Maynard HD;
 XX DR WPI; 2003-300420/29.
 XX PT Use of a ligand comprising of at least one sulfated or sulfonated amino
 PT acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic
 PT retinopathy and hypoxia.
 XX PS Disclosure; Fig 2; 79pp; English.
 XX CC The invention relates to a novel ligand for binding a target biomolecule,
 CC which comprises a peptide having at least one sulphated or sulphonated
 CC amino acid and at least one amino acid chosen from neutral and positively
 CC charged amino acids. The novel ligands can be used for the treatment of
 CC e.g. tumours, rheumatoid arthritis, diabetic retinopathy and hypoxia.
 CC This sequence represents a heparin binding peptide relating to the
 CC invention
 XX SQ Sequence 20 AA;

Query Match 76.1%; Score 33.5; DB 6; Length 20;
 Best Local Similarity 90.0%; Pred. No. 7.8;
 Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRK-GLK 9
 Db ||||| |||
 7 TRLTRKRLGLK 16

RESULT 15
 AAW57208
 ID AAW57208 standard; peptide; 22 AA.
 XX AC AAW57208;
 XX DT 03-AUG-1998 (first entry)
 XX DE Apo B 100 binding site peptide analogue peptide C.
 XX KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX OS Synthetic.
 XX FH Key Location/Qualifiers
 FT Modified-site 1 /note= "attached to retinoic acid"
 FT Modified-site 22 /note= "attached to cholesterol"
 XX PN WO9813385-A2.

XX 02-APR-1998.
XX PD
XX PF
XX 25-SEP-1997; 97WO-GB002610.
XX PR
XX 27-SEP-1996; 96GB-00020153.
XX PA (UYST) UNIV STRATHCLYDE.
XX PI Halbert GW, Owens MD, Baillie G;
XX WPI; 1998-230637/20.
XX Non-natural lipid particle comprising peptide binding to apo B protein
PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
PT that express this receptor.
XX Claim 13; Fig 7; 73pp; English.
XX The present sequence represents a specifically claimed Apo B 100 binding
CC site peptide analogue which can be used as a component of a non-
CC naturally occurring, receptor-competent low density lipoprotein (LDL)
CC particle of the present invention. The LDL particle comprises at least 1
CC peptide component that has at least 1 binding site for an apo B protein
CC receptor and at least 1 lipophilic substituent. Also described in the
CC invention are peptides containing an apo B binding sequence with at least
CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRGK (2), or their
CC dimers. Non-naturally occurring, receptor-competent LDL particles are
CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
CC cancer cells that express an apo B protein receptor, and (ii) additives
CC for cell culture media especially as growth supplements. Non-naturally
CC occurring, receptor-competent LDL particles do not require the complete
CC apo B sequence, which is large and tends to aggregate, to provide binding
CC affinity to an apo B protein receptor
XX SQ Sequence 22 AA;

Query Match 76.1%; Score 33.5; DB 2; Length 22;
Best Local Similarity 90.0%; Pred. No. 8.7;
Matches 9; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
Qy 1 TRLTRK-GLK 9
Db 7 TRLTRKRGK 16

Search completed: January 13, 2005, 01:43:03
Job time : 70.6585 secs

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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 70.2295 Seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-15
Perfect score: 44
Sequence: 1 TRLTRKGLK 9

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : UniProt_02: *
1: uniprot_sprot: *
2: uniprot_trembl: *

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB | ID | Description |
|------------|-------|-------------|--------|----|------------|-----------------------|
| 1 | 37 | 84.1 | 423 | 2 | Q7WSO9 | Q7wsq9 arthrobacte |
| 2 | 36 | 81.8 | 423 | 2 | O64892 | O64892 ananas como |
| 3 | 35 | 79.5 | 453 | 2 | O6BJK6 | O6bjk6 debaryomyce |
| 4 | 34 | 77.3 | 138 | 1 | R8FA_CHRVO | Q7ny12 chromobacte |
| 5 | 34 | 77.3 | 172 | 2 | Q9XBV5 | Q9xbv5 mycobacteri |
| 6 | 34 | 77.3 | 376 | 2 | Q7ZE67 | Q7ze67 desulfovibri |
| 7 | 34 | 77.3 | 376 | 2 | AA595192 | AA595192 desulfovibri |
| 8 | 34 | 77.3 | 451 | 1 | SYS_PYRAE | Q8ztp4 pyrobaculum |
| 9 | 34 | 77.3 | 457 | 2 | Q3FKN8 | Q3fkt8 arabidopsis |
| 10 | 34 | 77.3 | 482 | 2 | Q8Z5C5 | Q8z5c5 oryza sativ |
| 11 | 34 | 77.3 | 482 | 2 | BAD10124 | Bad10124 oryza sat |
| 12 | 34 | 77.3 | 493 | 2 | Q8TJF9 | Q8tjf9 methanosarc |
| 13 | 34 | 77.3 | 493 | 2 | Q8DNR6 | Q8dnr6 streptococc |
| 14 | 34 | 77.3 | 798 | 2 | Q97PA5 | Q97pa5 streptococc |
| 15 | 34 | 77.3 | 1230 | 2 | Q7G6A5 | Q7g6a5 oryza sativ |
| 16 | 34 | 77.3 | 1230 | 2 | Q8RU68 | Q8ru68 oryza sativ |
| 17 | 33.5 | 76.1 | 414 | 2 | Q7QR85 | Q7yqr5 aotus vocif |
| 18 | 33.5 | 76.1 | 596 | 2 | Q28473 | Q28473 macaca fasc |
| 19 | 33.5 | 76.1 | 3262 | 2 | Q13788 | Q13788 homo sapien |
| 20 | 33.5 | 76.1 | 4563 | 1 | APB_HUMAN | P04114 homo sapien |
| 21 | 33.5 | 76.1 | 4563 | 2 | Q7Z600 | Q7z600 homo sapien |
| 22 | 33 | 75.0 | 102 | 2 | Q8R2E8 | Q8r2e8 rattus norv |
| 23 | 33 | 75.0 | 137 | 2 | Q7MI23 | Q7mi23 vibrio vuln |
| 24 | 33 | 75.0 | 137 | 2 | Q8DBU7 | Q8dbu7 vibrio vuln |
| 25 | 33 | 75.0 | 147 | 1 | NDK_LISMO | Q8y5x4 listeria mo |
| 26 | 33 | 75.0 | 147 | 2 | Q7LY86 | Q7ly86 listeria mo |
| 27 | 33 | 75.0 | 147 | 2 | AA04728 | AA04728 listeria |
| 28 | 33 | 75.0 | 150 | 1 | RS13_BRUPA | P62299 brugia paha |
| 29 | 33 | 75.0 | 150 | 1 | RS13_WUCBA | P62300 wucheria |
| 30 | 33 | 75.0 | 151 | 1 | NDK_ARCFU | Q29491 archaeoglob |
| 31 | 33 | 75.0 | 151 | 2 | Q17274 | Q17274 brugia paha |

| | | | | | | |
|----|----|------|-----|---|------------|--------------------|
| 32 | 33 | 75.0 | 154 | 1 | NDK_METKA | Q8tv10 methanopyru |
| 33 | 33 | 75.0 | 157 | 2 | O9L533 | Q9l533 vibrio chol |
| 34 | 33 | 75.0 | 174 | 2 | O9KK25 | Q9kk25 vibrio chol |
| 35 | 33 | 75.0 | 306 | 2 | O87W29 | O87w29 pseudomonas |
| 36 | 33 | 75.0 | 335 | 2 | Q8PNK9 | Q8pnk9 xanthomonas |
| 37 | 33 | 75.0 | 470 | 1 | ROCC_BACSU | P39636 bacillus su |
| 38 | 33 | 75.0 | 503 | 2 | Q7P2I3 | Q7p2i3 fusobacteri |
| 39 | 33 | 75.0 | 506 | 2 | Q8REI1 | Q8rei1 fusobacteri |
| 40 | 33 | 75.0 | 592 | 2 | O6PAD4 | O6pap4 mus musculu |
| 41 | 33 | 75.0 | 592 | 2 | AAH57340 | AAh57340 mus muscu |
| 42 | 33 | 75.0 | 592 | 2 | AAH60177 | AAh60177 mus muscu |
| 43 | 33 | 75.0 | 716 | 2 | Q80YS3 | Q80ys3 mus musculu |
| 44 | 33 | 75.0 | 761 | 1 | PHT1_MOUSE | Q9qz09 mus musculu |
| 45 | 33 | 75.0 | 762 | 1 | PHT1_HUMAN | Q9ums5 homo sapien |

ALIGNMENTS

RESULT 1

Q7WSO9 PRELIMINARY; PRT; 423 AA.
AC Q7WSO9, PRELIMINARY; PRT; 423 AA.
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Putative transporter protein.
OS Arthrobacter ilicis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Micrococcineae; Micrococcaceae; Arthrobacter.
OX NCBI_TaxID=43665;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Rue61a;
RX MEDLINE=42753791; PubMed=12730200;
RA Parschat K., Hauer B., Kappel R., Kraft R., Huettermann J., Fetzner S.;
RT "Gene Cluster of Arthrobacter ilicis R.61a Involved in the Degradation
of Quinaldine to Anthranilate. Characterization and Functional
Expression of the Quinaldine 4-oxidase qoxLMS Genes.";
RL J. Biol. Chem. 278:27483-27494(2003).
DR EMBL; AJ537472; CAB61041.1; -.
DR GO; GO:0016021; C:integral to membrane; IEA.
DR GO; GO:0005215; P:transporter activity; IEA.
DR GO; GO:0006810; P:transport; IEA.
DR InterPro; IPR007114; MFS.
DR PROSITE; PSS0850; MFS; 1.
SQ SEQUENCE 423 AA; 43696 MW; BB11CBADA85DF241 CRC64;

Query Match 84.1%; Score 37; DB 2; Length 423;
Best Local Similarity 77.8%; Pred. No. 19;
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TRLTRKGLK 9

||||:||||
207 TRLTKQGLK 215

RESULT 2

O64892 PRELIMINARY; PRT; 871 AA.
ID O64892, PRELIMINARY; PRT; 871 AA.
AC O64892;
DT 01-AUG-1998 (TrEMBLrel. 07, Created)
DT 01-AUG-1998 (TrEMBLrel. 07, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Polyprotein (Fragment).
OS Ananas comosus (Pineapple).
OC Eukaryota; Viridiplantae; Streptophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Bromeliaceae;
OC Ananas.
OX NCBI_TaxID=4615;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=98418625; PubMed=9747853;

Thomson K.G., Thomas J.E., Dietzgen R.G.;
 "Retrotransposon-like sequences integrated into the genome of
 RT pineapple, *Ananas comosus*.";
 RL Plant Mol. Biol. 38:461-465(1998).
 DR EMBL: Y12432; CAA73042.1; -.
 DR PIR: T07863; T07863.
 DR GO: GO:0003677; F:DNA binding; IEA.
 DR GO: GO:0003723; F:RNA binding; IEA.
 DR GO: GO:0003964; F:RNA-directed DNA polymerase activity; IEA.
 DR GO: GO:0016740; F:transferase activity; IEA.
 DR GO: GO:0006310; P:DNA recombination; IEA.
 DR GO: GO:0006278; P:DNA-dependent DNA replication; IEA.
 DR InterPro: IPR001584; Rve.
 DR InterPro: IPR00477; RVTse.
 DR Pfam: PF00665; rve; 1.
 DR Pfam: PF00078; RVT; 1.
 KW Polyprotein; RNA-directed DNA polymerase; Transferase.
 FT NON_TER 871 871
 SQ SEQUENCE 871 AA; 100048 MW; EDFFD016E08952PC CRC64;
 Query Match 81.8%; Score 36; DB 2; Length 871;
 Best Local Similarity 77.8%; Pred. No. 72;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TRLTRKGLK 9
 DB 272 TRLTHKGVK 280
 RESULT 3
 Q6BJK6 PRELIMINARY; PRT; 453 AA.
 ID Q6BJK6
 AC Q6BJK6
 DT 01-OCT-2004 (TrEMBLrel. 28, Created)
 DT 01-OCT-2004 (TrEMBLrel. 28, Last sequence update)
 DT 01-OCT-2004 (TrEMBLrel. 28, Last annotation update)
 DE Similar to CA1599|IPF11452 *Candida albicans* IPF11452 of unknown
 DE function.
 GN ORFNames=DEHA0G02068g;
 OS Debaryomyces hansenii (Yeast) (Torulaspora hansenii).
 OC Eukaryota; Fungi; Ascomycota; Saccharomycotina; Saccharomycetes;
 OC Saccharomycetales; Saccharomycetaceae; Debaryomycetes.
 OX NCBI_TaxID=4959;
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=CRS767;
 RG GENOLEVURES;
 RA Dujon B., Sherman D., Fischer G., Durrens P., Casaregola S.,
 RA Lafontaine I., de Montigny J., Marck C., Neuvéglise C., Talla E.,
 RA Goffard N., Frangeul L., Aigle M., Anthouard V., Babour A., Barbe V.,
 RA Barnay S., Blanchin S., Beckerich J.M., Beyne E., Bleykasten C.,
 RA Boisarame A., Boyer J., Cattolico L., Confanioli F., de Daruvar A.,
 RA Despons L., Fabre E., Fairhead C., Ferry-Dumazet H., Groppi A.,
 RA Hantraye F., Hennequin C., Jauniaux N., Joyet P., Kachouri R.,
 RA Kerrest A., Koszul R., Lemaire M., Lesur I., Ma L., Muller H.,
 RA Nicaud J.M., Nikolski M., Oztas G., Ozier-Kalogeropoulos O.,
 RA Pellenz S., Potier S., Richard G.F., Straub M.L., Suleau A.,
 RA Swennene D., Tekia F., Wesolowski-Louvel M., Westhof E., Wirth B.,
 RA Zénou-Meyer M., Zivanovic I., Solotkin-Fukuhara M., Thierry A.,
 RA Bouchier P., Souciet J.L.;
 RT "Genome evolution in yeasts";
 RL Nature 430:35-44(2004).
 [2]
 RN SEQUENCE FROM N.A.
 RC STRAIN=CRS767;
 RA Genoscope;
 RL Submitted (JUL-2004) to the EMBL/GenBank/DBJ databases.
 DR EMBL: CR382139; CAG90062.1; -.
 SQ SEQUENCE 453 AA; 50757 MW; DD4AF51CF5D956F5 CRC64;
 Query Match 79.5%; Score 35; DB 2; Length 453;
 Best Local Similarity 77.8%; Pred. No. 58;
 Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
 QY 1 TRLTRKGLK 9
 DB 368 SRLTRKGTK 376
 RESULT 4
 RBFA_CHRVO STANDARD; PRT; 138 AA.
 ID Q7NY12;
 AC Q7NY12;
 DT 29-MAR-2004 (Rel. 43, Created)
 DT 29-MAR-2004 (Rel. 43, Last sequence update)
 DT 05-JUL-2004 (Rel. 44, Last annotation update)
 DE Ribosome-binding factor A.
 DE Names=rbfa; OrderedLocNames=CV1463;
 GN Chromobacterium violaceum.
 OS Bacteria; Proteobacteria; Betaproteobacteria; Neisseriales;
 OC Neisseriaceae; Chromobacterium.
 OX NCBI_TaxID=536;
 [1]
 RN SEQUENCE FROM N.A.
 RC STRAIN=ATCC 12472 / DSM 30191;
 RX MEDLINE=22882880; PubMed=14500782; DOI=10.1073/pnas.1832124100;
 RA Vasconcelos A.T.R., de Almeida D.F., Hungria M., Guimarães C.T.,
 RA Antonio R.V., Almeida F.C., de Almeida L.G.P., de Almeida R.,
 RA Alves-Gomes J.A., Andrade E.M., Araripe J., de Araujo M.F.F.,
 RA Astolfi-Filho S., Azevedo V., Baptista A.J., Bataus L.A.M.,
 RA Batista J.S., Belo A., van den Berg C., Bogo M., Bonatto S.,
 RA Bordignon J., Brígido M.M., Brito C.A., Brocchi M., Buriti H.A.,
 RA Camargo A.A., Cardoso D.D.P., Carneiro N.P., Carraro D.M.,
 RA Carvalho C.M.B., Cascado J.C.M., Cavada B.S., Chueire L.M.O.,
 RA Creczynski-Pasek T.B., Cunha-Junior N.C., Fagundes N., Faicao C.L.,
 RA Fancinatti F., Farias I.P., Felipe M.S.S., Fertari L.P., Ferro J.A.,
 RA Ferro M.I.T., Franco G.R., Freitas N.S.A., Furian L.R.,
 RA Gazzinelli R.T., Gomes E.A., Gonçalves P.R., Grangeiro T.B.,
 RA Grattapaglia D., Grigard E.C., Hanna E.S., Jardim S.N., Laurino J.,
 RA Leão L.C.T., Lima L.F.A., Loureiro M.F., Lyra M.C.C.P.,
 RA Madeira H.M.F., Manfio G.P., Maranhão A.O., Martins W.S.,
 RA di Mauro S.M.Z., de Medeiros S.R.B., Meissner R.V., Moreira M.A.M.,
 RA Nascimento F.F., Nicolas M.F., Oliveira J.G., Oliveira S.C.,
 RA Paixão R.C., Parente J.A., Pedrosa F.O., Pena S.D.J., Pereira J.O.,
 RA Pereira M., Pinto L.S.R.C., Pinto L.S., Porto J.I.R., Porrich D.P.,
 RA Ramalho-Neto C.E., Reis A.M.M., Rigo L.U., Rondinelli E.,
 RA Santos E.B.P., Santos F.R., Schneider M.P.C., Seuneh H.N.,
 RA Silva A.M.R., da Silva A.L.C., Silva D.W., Silva R., Simoes I.C.,
 RA Simon D., Soares C.M.A., Soares R.B.A., Souza E.M., Souza K.R.L.,
 RA Souza R.C., Steffens M.B.R., Steindel M., Teixeira S.R., Urmenyi T.,
 RA Vettore A., Wasson R., Zaha A., Simpson A.J.G.;
 RA "The complete genome sequence of *Chromobacterium violaceum* reveals
 RT remarkable and exploitable bacterial adaptability";
 RT Proc. Natl. Acad. Sci. U.S.A. 100:11660-11665(2003).
 RL CC -!- FUNCTION: Associates with free 30S ribosomal subunits (but not
 CC with 30S subunits that are part of 70S ribosomes or polyosomes).
 CC Essential for efficient processing of 16S rRNA. May interact with
 CC the 5'terminal helix region of 16S rRNA (By similarity).
 CC -!- SUBCELLULAR LOCATION: Cytoplasmic (Potential).
 CC -!- SIMILARITY: Belongs to the rbfa family.
 CC
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 CC
 CC EMBL: AE016915; AAQ59138.2; -.
 DR HAVAP; MF 00003; -. 1.
 DR InterPro: IPR000238; Rib_bind_factA.
 DR Pfam: PF02033; RBFA; 1.
 DR ProDom: PD007327; Rib_bind_factA; 1.

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DR TIGRFAMs; TIGR00082; rbfA; 1.
DR PROSITE; PS01319; RbFA; 1.
KW Complete proteome; rRNA processing.
SQ SEQUENCE 138 AA; 15387 MW; E84750D86390272C CRC64;

Query Match 77.3%; Score 34; DB 1; Length 138;
Best Local Similarity 100.0%; Pred.No. 26;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LTRKGLK 9
Db 25 LTRKGLK 31

RESULT 5
Q9XBV5 PRELIMINARY; PRT; 172 AA.
AC Q9XBV5;
DT 01-NOV-1999 (TrEMBLrel. 12, Created)
DT 01-NOV-1999 (TrEMBLrel. 12, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Hypothetical protein.
OS Mycobacterium smegmatis.
OC Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC Corynebacterineae; Mycobacteriaceae; Mycobacterium.
OX NCBI_TaxID=1772;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=mc2 155;
RA Bardou F., Martinez R., Puech V., Bleiber G., Prod'hom G., Daffe M.,
RA Telenti A.;
RL Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.
KW EMBL; AF155062; AAD42390.1; -.
KW Hypothetical protein.
SQ SEQUENCE 172 AA; 18622 MW; C4DFA92D0182E682 CRC64;

Query Match 77.3%; Score 34; DB 2; Length 172;
Best Local Similarity 100.0%; Pred.No. 33;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LTRKGLK 9
Db 132 LTRKGLK 138

RESULT 6
Q72E67 PRELIMINARY; PRT; 376 AA.
AC Q72E67;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DE Amino acid ABC transporter, periplasmic-binding protein.
GN OrderedLocusNames=DVU0712;
OS Desulfovibrio vulgaris (strain Hildenborough / ATCC 29579 / NCIMB 8303).
OC Bacteria; Proteobacteria; Deltaproteobacteria; Desulfovibrionales;
OC Desulfovibrionaceae; Desulfovibrio.
OX NCBI_TaxID=882;
RN [1]
RP SEQUENCE FROM N.A.
RX PubMed=15077118; DOI=10.1038/nbt959;
RA Heidelberg J.F., Seshadri R., Haveman S.A., Hemme C.L., Paulsen I.T.,
RA Kolonay J.F., Eisen J.A., Ward N.L., Methe B.A., Brinkac L.M.,
RA Daugherty S.C., DeBoy R.T., Dodson R.J., Durkin A.S., Madupu R.,
RA Nelson W.C., Sullivan S.A., Fouts D.E., Haft D.H., Selengut J.,
RA Peterson J.D., Davidlsen T.M., Zafar N., Zhou L., Radune D.,
RA Dmitrov G., Hance M., Tran K., Khouri H.M., Gill J., Utterback T.R.,
RA Feldblyum T.V., Wall J.D., Voordouw G., Fraser C.M.;
RT "The genome sequence of the anaerobic, sulfate-reducing bacterium
Desulfovibrio vulgaris Hildenborough."
RL Nat. Biotechnol. 22:554-559(2004).
DR EMBL; AE017311; AAS95192.1; -.
SQ SEQUENCE 376 AA; 40203 MW; 14F0AFE97CC2B79B CRC64;

Query Match 77.3%; Score 34; DB 2; Length 376;
Best Local Similarity 87.5%; Pred.No. 80;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLTRKGLK 9
Db 130 RLTRKGLK 137

RESULT 7
AAS95192 PRELIMINARY; PRT; 376 AA.
AC AAS95192;
DT 27-APR-2004 (TrEMBLrel. 27, Created)
DT 27-APR-2004 (TrEMBLrel. 27, Last sequence update)
DT 11-MAY-2004 (TrEMBLrel. 27, Last annotation update)
DE Amino acid ABC transporter, periplasmic-binding protein.
GN Desulfovibrio vulgaris (strain Hildenborough / ATCC 29579 / NCIMB 8303).
OC Bacteria; Proteobacteria; Deltaproteobacteria; Desulfovibrionales;
OC Desulfovibrionaceae; Desulfovibrio.
OX NCBI_TaxID=882;
RN [1]
RP SEQUENCE FROM N.A.
RX PubMed=15077118;
RA Heidelberg J.F., Seshadri R., Haveman S.A., Hemme C.L., Paulsen I.T.,
RA Kolonay J.F., Eisen J.A., Ward N., Methe B.A., Brinkac L.M.,
RA Daugherty S.C., DeBoy R.T., Dodson R.J., Durkin A.S., Madupu R.,
RA Nelson W.C., Sullivan S.A., Fouts D.E., Haft D.H., Selengut J.,
RA Peterson J.D., Davidlsen T.M., Zafar N., Zhou L., Radune D.,
RA Dmitrov G., Hance M., Tran K., Khouri H.M., Gill J., Utterback T.R.,
RA Feldblyum T.V., Wall J.D., Voordouw G., Fraser C.M.;
RT "The genome sequence of the anaerobic, sulfate-reducing bacterium
Desulfovibrio vulgaris Hildenborough."
RL Nat. Biotechnol. 22:554-559(2004).
DR EMBL; AE017311; AAS95192.1; -.
SQ SEQUENCE 376 AA; 40203 MW; 14F0AFE97CC2B79B CRC64;

Query Match 77.3%; Score 34; DB 2; Length 376;
Best Local Similarity 87.5%; Pred.No. 80;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKGLK 9
Db 130 RLTRKGLK 137

RESULT 8
SYS_PYRAE STANDARD; PRT; 451 AA.
AC Q8ZTP4;
DT 10-OCT-2003 (Rel. 42, Created)
DT 10-OCT-2003 (Rel. 42, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Seryl-tRNA synthetase (EC 6.1.1.11) (Serine--tRNA ligase) (SerRS).
GN Name=serS; OrderedLocusNames=PAE1158;
OS Pyrobaculum aerophilum.
OC Archaea; Crenarchaeota; Thermoprotei; Thermoproteales;
OC Thermoproteaceae; Pyrobaculum.
OX NCBI_TaxID=13773;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=IM2 / ATCC 51768 / DSM 7523;

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RX PubMed=11792869; DOI=10.1073/pnas.241636498;
 RA Fitz-Gibbon S.T., Ladner H., Kim U.-J., Stetter K.O., Simon M.I.,
 RA Miller J.H.;
 RT "Genome sequence of the hyperthermophilic crenarchaeon Pyrobaculum
 aerophilum";
 RL Proc. Natl. Acad. Sci. U.S.A. 99:984-989 (2002).
 CC -|- CATALYTIC ACTIVITY: ATP + L-serine + tRNA(Ser) = AMP + diphosphate
 + L-seryl-tRNA(Ser).
 CC -|- SUBCELLULAR LOCATION: Cytoplasmic.
 CC -|- SIMILARITY: Belongs to class-II aminoacyl-tRNA synthetase family.
 CC
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 CC
 CC EMBL; AE009914; AAL64715.1; -;
 CC HSSP; P34945; 1SER.
 DR HAMAP; MF 00176; -; 1.
 DR InterPro; IPR002314; tRNA-synt_2b.
 DR InterPro; IPR002317; tRNA-synt_ser.
 DR InterPro; IPR010978; tRNA binding arm.
 DR InterPro; IPR006195; tRNA ligase_II.
 DR Pfam; PF02403; Seryl tRNA N; 1.
 DR Pfam; PF00587; tRNA-synt_2b; 1.
 DR PRINTS; PR00981; TRNASYNTHSER.
 DR TIGRFAMs; TIGR00414; sers; 1.
 DR PROSITE; PS50862; AA TRNA LIGASE_II; 1.
 KW Aminoacyl-tRNA synthetase; ATP-binding; Complete proteome; Ligase;
 KW Protein biosynthesis.
 SQ SEQUENCE 451 AA; 52030 MW; 985B4C826E75D505 CRC64;

Query Match 77.3%; Score 34; DB 1; Length 451;
 Best Local Similarity 75.0%; Pred. No. 98;
 Matches 6; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKGLK 9
 |:|||||:
 Db 383 RVRKGMK 390

RESULT 9
 Q9FKT8 PRELIMINARY; PRT; 457 AA.
 AC Q9FKT8;
 DT 01-MAR-2001 (TrEMBLrel. 16, Created)
 DT 01-MAR-2001 (TrEMBLrel. 16, Last sequence update)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
 DE Similarity to cytochrome oxidase assembly factor (Hypothetical protein
 At5g56090/MDA7 15) (Hypothetical protein At5g56090).
 GN Name=At5g56090/MDA7 15; Synonyms=At5g56090;
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
 OC eurosids II; Brassicales; Brassicaceae; Arabidopsi.
 OX NCBI_TaxID=3702;
 [1]
 RP SEQUENCE FROM N.A.
 RA MEDLINE=98344145; PubMed=9679202;
 RA Kaneko T., Kotani H., Nakamura Y., Sato S., Asamizu E., Miyajima N.,
 RA Tabata S.;
 RT "Structural analysis of Arabidopsis thaliana chromosome 5. V. Sequence
 RT features of the regions of 1,381,565 bp covered by twenty one
 RT physically assigned P1 and TAC clones.";
 RL DNA Res. 5:131-145 (1998).
 RN
 RN SEQUENCE FROM N.A.
 RA Seki M., Iida K., Satou M., Sakurai T., Akiyama K., Ishida J.,
 RA Nakajima M., Enju A., Kamiya A., Narusaka M., Carninci P., Kawai J.,
 RA Hayaishizaki Y., Shinozaki K.

RL Submitted (NOV-2002) to the EMBL/GenBank/DBJ databases.
 RN [3]

RP SEQUENCE FROM N.A.
 RA Yamada K., Chan M.M., Chang C.H., Dale J.M., Hsuan V.W., Lee J.M.,
 RA Onodera C.S., Quach H.L., Tang C., Toriumi M., Wong C., Wu H.C.,
 Yu G., Yuan S., Carninci P., Chen H., Cheuk R., Hayashizaki Y.,
 RA Ishida J., Jones T., Kamiya A., Kawai J., Kim C.J., Narusaka M.,
 RA Nguyen M., Palm C.J., Sakurai T., Satou M., Seki M., Shinn P.,
 RA Southwick A., Tripp M.G., Wu T., Shinozaki K., Davis R.W., Ecker J.R.,
 RA Theologis A.;
 RL Submitted (MAR-2003) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AB011476; BAB09291.1; -;
 DR EMBL; AK117496; BAC42159.1; -;
 DR EMBL; BT004976; AAC050509.1; -;
 DR GO; GO:001620; C:membrane; IEA.
 DR GO; GO:0006461; P:protein complex assembly; IEA.
 DR InterPro; IPR003780; COX15_CtaA.
 DR Pfam; PF02628; COX15_CtaA; 1.
 KW Hypothetical protein.
 SQ SEQUENCE 457 AA; 50127 MW; 6E7A0DE6457E2D1E CRC64;

Query Match 77.3%; Score 34; DB 2; Length 457;
 Best Local Similarity 87.5%; Pred. No. 99;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TLTRKGL 8
 |:|||||:
 Db 120 TLTRSGL 127

RESULT 10
 Q6ZSC5 PRELIMINARY; PRT; 482 AA.
 ID Q6ZSC5;
 AC Q6ZSC5;
 DT 05-JUL-2004 (TrEMBLrel. 27, Created)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
 DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
 DE Putative cytochrome c oxidase subunit 15(COX15) homolog isoform
 DE 1.
 GN Name=B1142B04.27;
 OS Oryza sativa (japonica cultivar-group).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
 OC Ehrhartoideae; Oryzeae; Oryza.
 OX NCBI_TaxID=39947;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Sasaki T., Matsumoto T., Katayose Y.;
 RL Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
 DR EMBL; AP005148; BAD10124.1; -;
 DR InterPro; IPR003780; COX15_CtaA.
 DR Pfam; PF02628; COX15_CtaA; 1.
 SQ SEQUENCE 482 AA; 51446 MW; DC46727EC38CEAD5 CRC64;

Query Match 77.3%; Score 34; DB 2; Length 482;
 Best Local Similarity 87.5%; Pred. No. 1.1e+02;
 Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TLTRKGL 8
 |:|||||:
 Db 147 TLTRSGL 154

RESULT 11
 BAD10124 PRELIMINARY; PRT; 482 AA.
 ID BAD10124;
 AC BAD10124;
 DT 02-MAR-2004 (TrEMBLrel. 27, Created)
 DT 02-MAR-2004 (TrEMBLrel. 27, Last sequence update)
 DT 02-MAR-2004 (TrEMBLrel. 27, Last annotation update)
 DE Putative cytochrome c oxidase subunit 15(COX15) homolog isoform
 DE 1.
 GN B1142B04.27.

```

OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzaceae; Oryza; Oryza sativa.
OX NCBI_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=cv. Nipponbare;
RA Sasaki T., Matsumoto T., Katayose Y.;
RT "Oryza sativa nipponbare (GA3) genomic DNA, chromosome 8, BAC
clone:B1142B04.";
RL Submitted (MAY-2002) to the EMBL/GenBank/DBSJ databases.
DR EMBL; AP005148; BAD10124.1; -.
SQ SEQUENCE 482 AA; 51446 MW; DC46727EC38CEAD5 CRC64;

Query Match
Best Local Similarity 77.3%; Score 34; DB 2; Length 482;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKGL 8
    |||||
Db 147 TRLTRSGL 154

RESULT 12
Q8TJF9 PRELIMINARY; PRT; 493 AA.
AC Q8TJF9;
DT 01-JUN-2002 (TREMBlrel. 21, Created)
DT 01-JUN-2002 (TREMBlrel. 21, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Terminase.
GN OrderedLocusNames=MA3826;
OS Methanosarcina acetivorans.
OC Archaea; Euryarchaeota; Methanomicrobia; Methanosarcinales;
OC Methanosarcinaceae; Methanosarcina.
OX NCBI_TaxID=2214;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=C2A / ATCC 35395 / DSM 2834;
EX MEDLINE=2129760; PubMed=1932238; DOI=10.1101/gr.223902;
RA Galagan J.E., Nussbaum C., Roy A., Endrizzi M.G., Macdonald P.,
RA FitzHugh W., Calvo S., Engels R., Smirnov S., Atnoor D., Brown A.,
RA Allen N., Naylor J., Stange-Thomann N., DeArellano K., Johnson R.,
RA Linton L., McEwan P., McKernan K., Talamas J., Tirrell A., Ye W.,
RA Zimmer A., Barber R.D., Cann I., Graham D.E., Grahame D.A., Guss A.M.,
RA Hedderich R., Ingram-Smith C., Kuettnner H.C., Krzycki J.A., Smith K.,
RA Leigh J.A., Li W., Liu J., Mukhopadhyay B., Reeve J.N., Smith K.,
RA Springer T.A., Unayam L.A., White O., White R.H., de Macario E.C.,
RA Ferry J.G., Jarrell K.F., Jing H., Macario A.J.L., Paulsen I.T.,
RA Pritchett M., Sowers K.R., Swanson R.V., Zinder S.H., Lander E.,
RA Metcalf W.W., Birren B.;
RT "The genome of Methanosarcina acetivorans reveals extensive metabolic
and physiological diversity.";
RL Genome Res. 12:532-542(2002).
DR EMBL; AE011094; AA007177.1; -.
DR InterPro; IPR006517; DUF phage C.
DR TIGRFAMs; TIGR01630; psiM2_ORF3; 1.
KW Complete proteome.
SQ SEQUENCE 493 AA; 56992 MW; 0170B5011675FD0C CRC64;

Query Match
Best Local Similarity 100.0%; Score 34; DB 2; Length 493;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 3 LTRKGLK 9
    |||||
Db 473 LTRKGLK 479

RESULT 13
Q8DNR6 PRELIMINARY; PRT; 798 AA.
ID Q8DNR6

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AC Q8DNR6;
DT 01-MAR-2003 (TREMBlrel. 23, Created)
DT 01-MAR-2003 (TREMBlrel. 23, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Primosomal replication factor Y.
GN Name=prfA; OrderedLocusNames=sp1581;
OS Streptococcus pneumoniae (strain ATCC BAA-255 / R6).
OC Bacteria; Firmicutes; Lactobacillales; Streptococcaceae;
OC Streptococcus.
OX NCBI_TaxID=171101;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21429245; PubMed=11544234;
RA Hoekins J., Alborn W.E. Jr., Arnold J., Blaszcak L.C., Burgett S.,
RA Dehoff B.S., Estrem S.T., Fritz L., Fu D.-J., Fuller W., Geringer C.,
RA Gilmore R., Glass J.S., Khoja H., Kraft A.R., Lagace R.E.,
RA Leblanc D.J., Lee L.N., Lefkowitz E.J., Lu J., Matsushima P.,
RA McAhren S.M., McHenry M., Mcleaster K., Mundy C.W., Nickas T.I.,
RA Norris F.H., O'Gara M., Peery R.B., Robertson G.T., Rockey P.,
RA Sun P.-M., Winkler M.E., Yang Y., Young-Bellido M., Zhao G.,
RA Zook C.A., Baltz R.H., Jaskunas S.R., Rostock P.R. Jr., Skatrud P.L.,
RA Glass J.I.;
RT "Genome of the bacterium Streptococcus pneumoniae strain R6.";
RL J. Bacteriol. 183:5709-5717(2001).
DR EMBL; AE008525; AAL00384.1; -.
DR PIR; C98069; C98069.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0008026; F:ATP-dependent helicase activity; IEA.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR GO; GO:0003676; F:nucleic acid binding; IEA.
DR GO; GO:0015668; F:type III site-specific deoxyribonuclease ac. .; IEA.
DR GO; GO:0009307; P:DNA restriction; IEA.
DR InterPro; IPR001410; DEAD.
DR InterPro; IPR001650; Helicase_C.
DR InterPro; IPR005259; PrfA.
DR InterPro; IPR006935; ResIII.
DR Pfam; PF00271; Helicase_C; 1.
DR Pfam; PF04851; ResIII; 1.
DR SMART; SM00487; DEXDc; 1.
DR SMART; SM00490; HELICc; 1.
DR TIGRFAMs; TIGR00595; prfA; 1.
DR ATP-binding; Complete proteome; Helicase; Hydrolase.
SQ SEQUENCE 798 AA; 90036 MW; DDOC4EEA5B269962 CRC64;

Query Match
Best Local Similarity 100.0%; Score 34; DB 2; Length 798;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKGL 8
    |||||
Db 156 RLTRKGL 162

RESULT 14
Q97PA5 PRELIMINARY; PRT; 798 AA.
AC Q97PA5;
DT 01-OCT-2001 (TREMBlrel. 18, Created)
DT 01-OCT-2001 (TREMBlrel. 18, Last sequence update)
DT 01-OCT-2003 (TREMBlrel. 25, Last annotation update)
DE Primosomal protein N'.
GN OrderedLocusNames=SP1736;
OS Streptococcus pneumoniae.
OC Bacteria; Firmicutes; Lactobacillales; Streptococcaceae;
OC Streptococcus.
OX NCBI_TaxID=1313;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=ATCC BAA-334 / TIGR4;
RX MEDLINE=21357209; PubMed=11463916; DOI=10.1126/science.1061217;
RA Tettelin H., Nelson K.E., Paulsen I.T., Eisen J.A., Read T.D.,
RA Peterson S.N., Heidelberg J.F., DeBoy R.T., Haft D.H., Dodson R.J.,
RA Durkin A.S., Gwinn M.L., Kolonay J.F., Nelson W.C., Peterson J.D.,

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RA Umayam L.A., White O., Salzberg S.L., Lewis M.R., Radune D.,
RA Holtzaple E.K., Khouri H.M., Wolf A.M., Utterback T.R., Hansen C.L.,
RA McDonald L.A., Feldblyum T.V., Angiuoli S.V., Dickinson T.,
RA Hickey E.K., Holt J.E., Loftus B.J., Yang E., Smith H.O., Venter J.C.,
RA Dougherty B.A., Morrison D.A., Hollingshead S.K., Fraser C.M.;
RT "Complete genome sequence of a virulent isolate of Streptococcus
RT pneumoniae.";
RL Science 293:498-506(2001).
DR EMBL; AE007466; AAK75812.1; -.
DR PIR; C95202; C95202.
DR TIGR; SP1736; -.
DR GO; GO:0005524; F:ATP binding; IEA.
DR GO; GO:0008026; F:ATP-dependent helicase activity; IEA.
DR GO; GO:0016787; F:hydrolase activity; IEA.
DR GO; GO:0003676; F:nucleic acid binding; IEA.
DR InterPro; IPR001410; DEAD.
DR InterPro; IPR001650; Helicase_C.
DR InterPro; IPR005259; PriA.
DR Pfam; PF00271; Helicase_C; 1.
DR SMART; SM00487; DEXDC; 1.
DR SMART; SM00490; HELICG; 1.
DR TIGRFAMs; TIGR00595; priA; 1.
KW ATP-binding; Complete proteome; Helicase; Hydrolase.
SQ SEQUENCE 798 AA; 9028 MW; FED84D4A8BB9B198 CRC64;

Query Match 77.3%; Score 34; DB 2; Length 798;
Best Local Similarity 100.0%; Pred. No. 1.9e+02;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKGL 8
Db 156 RLTRKGL 162

RESULT 15
Q7G6A5 PRELIMINARY; PRT; 1230 AA.
AC Q7G6A5;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DE Putative 22 kDa kafirin cluster; Ty3-Gypsy type.
GN ORFNames=OSJNAB0075K12.33;
OS Oryza sativa (japonica cultivar-group).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
OC Ehrhartoideae; Oryzoae; Oryza.
OC NCB1_TaxID=39947;
RN [1]
RP SEQUENCE FROM N.A.
RA Wing R.A., Yu Y., Yang T.J., Nah G., Soderlund C., Chen M., Kim H.-R.,
RA Rambo T., Saeki C., Henry D., Oates R., Simmons J.;
RL Submitted (JUN-2002) to the EMBL/GenBank/DBJ databases.
RN [2]
RP SEQUENCE FROM N.A.
RA The Rice Chromosome 10 Sequencing Consortium;
RT "In-depth view of structure, activity, and evolution of rice
RT chromosome 10.";
RL Science 300:1566-1569(2003).
RN [3]
RP SEQUENCE FROM N.A.
RA Buell C.R., Wing R.A., McCombie W.R., Messing J., Yuan Q.;
RL Submitted (MAY-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AC122148; AAM48279.1; -.
DR EMBL; AE017082; AAP53268.1; -.
DR InterPro; IPR009007; Pept_Aspartic.
DR InterPro; IPR001969; Pept_Asp_AS.
DR InterPro; IPR005162; Retrotrans_gag.
DR InterPro; IPR001584; Rve.
DR InterPro; IPR0000477; RVTse.
DR InterPro; IPR001878; Znf_CCHC.
DR Pfam; PF03732; Retrotran_gag; 1.
DR Pfam; PF00665; rve; 1.

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DR Pfam; PF00078; RVT; 1.
DR Pfam; PF00098; zf_CCHC; 1.
DR PROSITE; PS00141; ASP_PROTEASE; UNKNOWN_1.
DR PROSITE; PSS0158; zf_CCHC; 1.
KW RNA-directed DNA polymerase; Transferase.
SQ SEQUENCE 1230 AA; 141401 MW; 28B6664530A45846 CRC64;

Query Match 77.3%; Score 34; DB 2; Length 1230;
Best Local Similarity 66.7%; Pred. No. 3e+02;
Matches 6; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TRLTQKGLK 9
Db 706 TRLTQKGLK 714

Search completed: January 13, 2005, 01:51:05
Job time : 72.2295 secs

```

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 84.9344 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-17
Perfect score: 54
Sequence: 1 TRLTRKERGLK 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : A_Geneseq_23Sep04.*
1: Geneseqp1980s.*
2: Geneseqp1990s.*
3: Geneseqp2000s.*
4: Geneseqp2001s.*
5: Geneseqp2002s.*
6: Geneseqp2003as.*
7: Geneseqp2003bs.*
8: Geneseqp2004s.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB | ID | Description |
|------------|-------|-------------|--------|----|----------|-------------|
| 1 | 54 | 100.0 | 11 | 2 | AA30698 | Apo-B100 |
| 2 | 51 | 94.4 | 11 | 2 | AA30700 | Apo-B100 |
| 3 | 46 | 85.2 | 11 | 2 | AA30697 | Apo-B100 |
| 4 | 44 | 81.5 | 11 | 2 | AA30699 | Apo-B100 |
| 5 | 38.5 | 71.3 | 10 | 2 | AA30690 | Apo-B100 |
| 6 | 38.5 | 71.3 | 10 | 2 | AA30682 | Apo-B100 |
| 7 | 38.5 | 71.3 | 11 | 2 | AAW57205 | Apo B bin |
| 8 | 38.5 | 71.3 | 13 | 2 | AAW57207 | Apo B 100 |
| 9 | 38.5 | 71.3 | 15 | 2 | AAW41261 | Apolipop |
| 10 | 38.5 | 71.3 | 15 | 2 | AAW96892 | Apob-100 |
| 11 | 38.5 | 71.3 | 20 | 6 | ABJ37575 | Heparin b |
| 12 | 38.5 | 71.3 | 22 | 2 | AAW57208 | Apo B 100 |
| 13 | 38.5 | 71.3 | 22 | 2 | AAW57209 | Apo B 100 |
| 14 | 38.5 | 71.3 | 34 | 5 | AAE14541 | Human apo |
| 15 | 38.5 | 71.3 | 36 | 2 | AAW96876 | Nucleic a |
| 16 | 38.5 | 71.3 | 37 | 2 | AAW64587 | Human apo |
| 17 | 38.5 | 71.3 | 51 | 2 | AAW96845 | Nucleic a |
| 18 | 38.5 | 71.3 | 343 | 4 | ABB37687 | Peptide # |
| 19 | 38.5 | 71.3 | 343 | 4 | ABG52504 | Human liv |
| 20 | 38.5 | 71.3 | 377 | 2 | AAW72704 | Human apo |
| 21 | 38.5 | 71.3 | 377 | 2 | AAW34031 | Sequence |
| 22 | 38.5 | 71.3 | 2463 | 8 | ADJ57400 | Human apo |
| 23 | 38.5 | 71.3 | 3923 | 2 | AAW31237 | Human apo |
| 24 | 38.5 | 71.3 | 4536 | 2 | AAW41262 | Apolipop |
| 25 | 38.5 | 71.3 | 4536 | 2 | AAW96826 | Amino aci |

| | | | | | | |
|----|------|------|------|---|----------|-----------|
| 26 | 38.5 | 71.3 | 4560 | 5 | AAU98981 | Human apo |
| 27 | 38.5 | 71.3 | 4561 | 7 | ADD48677 | Human pro |
| 28 | 38.5 | 71.3 | 4563 | 5 | AAO15893 | Human apo |
| 29 | 38.5 | 71.3 | 4563 | 6 | ABR40253 | Human ali |
| 30 | 38.5 | 71.3 | 4563 | 6 | ABU79140 | Apolipop |
| 31 | 38.5 | 71.3 | 4563 | 7 | ADF43408 | Apolipop |
| 32 | 38.5 | 71.3 | 4563 | 8 | ADH18871 | Human apo |
| 33 | 38.5 | 71.3 | 4563 | 8 | ADH18870 | Human apo |
| 34 | 38.5 | 71.3 | 4563 | 8 | ADO33445 | Human apo |
| 35 | 38.5 | 71.3 | 4563 | 8 | ADO33447 | Human apo |
| 36 | 38.5 | 71.3 | 4590 | 4 | AAU33184 | Novel hum |
| 37 | 38 | 70.4 | 97 | 4 | AAU21930 | Human car |
| 38 | 38 | 70.4 | 97 | 7 | ADE45898 | Human car |
| 39 | 38 | 70.4 | 270 | 4 | ABG14088 | Novel hum |
| 40 | 38 | 70.4 | 1023 | 4 | ABG06061 | Novel hum |
| 41 | 37 | 68.5 | 488 | 5 | ABB93444 | Herbicida |
| 42 | 36 | 66.7 | 1095 | 5 | ABP52170 | Mouse pot |
| 43 | 36 | 66.7 | 1095 | 7 | ADD37509 | Mouse pho |
| 44 | 36 | 66.7 | 1095 | 8 | ADI27988 | Murine pr |
| 45 | 35.5 | 65.7 | 10 | 2 | AA30683 | Apo-B100 |

ALIGNMENTS

RESULT 1
AA30698
ID AA30698 standard; peptide; 11 AA.
XX AC AA30698;
XX 17-NOV-1999 (first entry)
XX Apo-B100 derived peptide showing a proteoglycan receptor mutation.
DE Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX Synthetic.
OS Homo sapiens.
XX WO9946598-A1.
XX 16-SEP-1999.
XX 05-MAR-1999; 99WO-US004805.
XX 10-MAR-1998; 98US-0077618P.
XX (REGC) UNIV CALIFORNIA.
XX Innerarity TL, Boren JOS;
XX WPI; 1999-551509/46.
XX Identifying compounds which affect binding of low density lipoprotein
XX with proteoglycan, used for, e.g. obtaining compounds for reducing
XX atherosclerosis.
XX Claim 17; Page 57; 70pp; English.

AA30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
receptor mutations. They were created to identify compounds which
modulate atherosclerosis. The peptides are derived from amino acids 3358
to 3367 of apoB100. The method comprises detecting compounds which affect
low density lipoprotein (LDL) binding with proteoglycan (PG). The method
can be used for identifying compounds which disrupt LDL-PG binding
without inhibiting LDL receptor binding. Such compounds can be used to
reduce or prevent the formation of atherosclerotic lesions and prevent
atherosclerosis. The transgenic non-human animals and mammals which
express human apo-B100 can be used as an in vivo model system for the
study of atherosclerosis, and in vivo assay methods for identifying
compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal

XX SQ Sequence 11 AA;

Query Match 100.0%; Score 54; DB 2; Length 11;
 Best Local Similarity 100.0%; Pred. No. 0.0044;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKERGLK 11
 |||||
 Db 1 TRLTRKERGLK 11

RESULT 2

AAV30700
 ID AAY30700 standard; peptide; 11 AA.

XX AC AAY30700;

XX DT 17-NOV-1999 (first entry)

XX DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

XX KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

XX OS Synthetic.

XX OS Homo sapiens.

XX PN WO9946598-A1.

XX PD 16-SEP-1999.

XX PF 05-MAR-1999; 99WO-US004805.

XX PR 10-MAR-1998; 98US-0077618P.

XX PA (REGC) UNIV CALIFORNIA.

XX PI Innerarity TL, Boren JOS;

XX DR WPI; 1999-551509/46.

XX CC Identifying compounds which affect binding of low density lipoprotein
 with proteoglycan, used for, e.g. obtaining compounds for reducing
 atherosclerosis.

XX PS Claim 17; Page 58; 70pp; English.

XX CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 receptor mutations. They were created to identify compounds which
 modulate atherosclerosis. The peptides are derived from amino acids 3358
 to 3367 of apoB100. The method comprises detecting compounds which affect
 low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 can be used for identifying compounds which disrupt LDL-PG binding
 without inhibiting LDL receptor binding. Such compounds can be used to
 reduce or prevent the formation of atherosclerotic lesions and prevent
 atherosclerosis. The transgenic non-human animals and mammals which
 express human apo-B100 can be used as an in vivo model system for the
 study of atherosclerosis, and in vivo assay methods for identifying
 compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 also be used to identify compounds which result in an increase in
 atherosclerotic regions. Thus the assays may be used to determine whether
 a particular food or drug composition tends to stimulate or inhibit the
 formation of atherosclerotic lesions. The polynucleotides can also be
 used in gene therapy for preventing or reducing the severity of
 atherosclerosis in an animal or mammal

XX

SQ Sequence 11 AA;

Query Match 94.4%; Score 51; DB 2; Length 11;
 Best Local Similarity 90.9%; Pred. No. 0.016;
 Matches 10; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKERGLK 11
 |||||
 Db 1 TRLTRKERGLK 11

RESULT 3

AAV30697
 ID AAY30697 standard; peptide; 11 AA.

XX AC AAY30697;

XX DT 17-NOV-1999 (first entry)

XX DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

XX KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

XX OS Synthetic.

XX OS Homo sapiens.

XX PN WO9946598-A1.

XX PD 16-SEP-1999.

XX PF 05-MAR-1999; 99WO-US004805.

XX PR 10-MAR-1998; 98US-0077618P.

XX PA (REGC) UNIV CALIFORNIA.

XX PI Innerarity TL, Boren JOS;

XX DR WPI; 1999-551509/46.

XX CC Identifying compounds which affect binding of low density lipoprotein
 with proteoglycan, used for, e.g. obtaining compounds for reducing
 atherosclerosis.

XX PS Claim 17; Page 57; 70pp; English.

XX CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 receptor mutations. They were created to identify compounds which
 modulate atherosclerosis. The peptides are derived from amino acids 3358
 to 3367 of apoB100. The method comprises detecting compounds which affect
 low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 can be used for identifying compounds which disrupt LDL-PG binding
 without inhibiting LDL receptor binding. Such compounds can be used to
 reduce or prevent the formation of atherosclerotic lesions and prevent
 atherosclerosis. The transgenic non-human animals and mammals which
 express human apo-B100 can be used as an in vivo model system for the
 study of atherosclerosis, and in vivo assay methods for identifying
 compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 also be used to identify compounds which result in an increase in
 atherosclerotic regions. Thus the assays may be used to determine whether
 a particular food or drug composition tends to stimulate or inhibit the
 formation of atherosclerotic lesions. The polynucleotides can also be
 used in gene therapy for preventing or reducing the severity of
 atherosclerosis in an animal or mammal

SQ Sequence 11 AA;

Query Match 85.2%; Score 46; DB 2; Length 11;
 Best Local Similarity 81.8%; Pred. No. 0.13;
 Matches 9; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKERGLK 11


```

Db      1  TRLTRKRGK 11
|||||:||||
RESULT 4
RAY30699
ID  AAY30699 standard, peptide; 11 AA.
XX
AC  AAY30699;
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW  Apo-B100; proteoglycan receptor mutation; atherosclerosis;
XX  low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS  Synthetic.
XX  Homo sapiens.
XX
PN  WO9946598-A1.
XX
PD  16-SEP-1999.
XX
PF  05-MAR-1999; 99WO-US004805.
XX
PR  10-MAR-1998; 98US-0077618P.
XX
PA  (REGC ) UNIV CALIFORNIA.
XX
PI  Innerarity TL, Boren JOS;
XX
DR  WPI; 1999-551509/46.
XX
PT  Identifying compounds which affect binding of low density lipoprotein
PT  with proteoglycan, used for, e.g. obtaining compounds for reducing
PT  atherosclerosis.
XX
PS  Claim 17; Page 58; 70pp; English.
XX
CC  AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC  receptor mutations. They were created to identify compounds which
CC  modulate atherosclerosis. The peptides are derived from amino acids 3358
CC  to 3367 of apoB100. The method comprises detecting compounds which affect
CC  low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC  can be used for identifying compounds which disrupt LDL-PG binding
CC  without inhibiting LDL receptor binding. Such compounds can be used to
CC  reduce or prevent the formation of atherosclerotic lesions and prevent
CC  atherosclerosis. The transgenic non-human animals and mammals which
CC  express human apo-B100 can be used as an in vivo model system for the
CC  study of atherosclerosis, and in vivo assay methods for identifying
CC  compounds which modulate atherosclerosis and/or LDL-PG binding. They can
CC  also be used to identify compounds which result in an increase in
CC  atherosclerotic regions. Thus the assays may be used to determine whether
CC  a particular food or drug composition tends to stimulate or inhibit the
CC  formation of atherosclerotic lesions. The polynucleotides can also be
CC  used in gene therapy for preventing or reducing the severity of
CC  atherosclerosis in an animal or mammal
XX
SQ  Sequence 11 AA;
Query Match 81.5%; Score 44; DB 2; Length 11;
Best Local Similarity 81.8%; Pred. No. 0.3;
Matches 9; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy      1  TRLTRKRGK 11
|||||:||||
Db      1  TRLTRKRGK 11
|||||:||||

RESULT 5
AAY30690
ID  AAY30690 standard, peptide; 10 AA.
XX
AC  AAY30690;
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW  Apo-B100; proteoglycan receptor mutation; atherosclerosis;
XX  low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS  Synthetic.
XX  Homo sapiens.
XX
PN  WO9946598-A1.
XX
PD  16-SEP-1999.
XX
PF  05-MAR-1999; 99WO-US004805.
XX
PR  10-MAR-1998; 98US-0077618P.
XX
PA  (REGC ) UNIV CALIFORNIA.
XX
PI  Innerarity TL, Boren JOS;
XX
DR  WPI; 1999-551509/46.
XX
PT  Identifying compounds which affect binding of low density lipoprotein
PT  with proteoglycan, used for, e.g. obtaining compounds for reducing
PT  atherosclerosis.
XX
PS  Claim 17; Page 57; 70pp; English.
XX
CC  AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC  receptor mutations. They were created to identify compounds which
CC  modulate atherosclerosis. The peptides are derived from amino acids 3358
CC  to 3367 of apoB100. The method comprises detecting compounds which affect
CC  low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC  can be used for identifying compounds which disrupt LDL-PG binding
CC  without inhibiting LDL receptor binding. Such compounds can be used to
CC  reduce or prevent the formation of atherosclerotic lesions and prevent
CC  atherosclerosis. The transgenic non-human animals and mammals which
CC  express human apo-B100 can be used as an in vivo model system for the
CC  study of atherosclerosis, and in vivo assay methods for identifying
CC  compounds which modulate atherosclerosis and/or LDL-PG binding. They can
CC  also be used to identify compounds which result in an increase in
CC  atherosclerotic regions. Thus the assays may be used to determine whether
CC  a particular food or drug composition tends to stimulate or inhibit the
CC  formation of atherosclerotic lesions. The polynucleotides can also be
CC  used in gene therapy for preventing or reducing the severity of
CC  atherosclerosis in an animal or mammal
XX
SQ  Sequence 10 AA;
Query Match 71.3%; Score 38.5; DB 2; Length 10;
Best Local Similarity 90.9%; Pred. No. 2.8;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy      1  TRLTRKRGK 11
|||||:||||
Db      1  TRLTRKE-GLK 10
|||||:||||

RESULT 6
AAY30682
ID  AAY30682 standard; peptide; 10 AA.
XX
AC  AAY30682;
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX

```


XX (UYST) UNIV STRATHCLYDE.
 XX Halbert GW, Owens MD, Baillie G;
 XX WPI; 1998-230637/20.
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX Claim 13; Fig 7; 73pp; English.
 XX The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRGK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 13 AA;
 Query Match 71.3%; Score 38.5; DB 2; Length 13;
 Best Local Similarity 90.9%; Pred. No. 3.6;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRKRGK 11
 Db 3 TRLTRK-RGLK 12
 RESULT 9
 AAW41261
 ID AAW41261 standard; peptide; 15 AA.
 XX
 AC AAW41261;
 XX
 DT 19-MAY-1998 (first entry)
 XX
 DE Apolipoprotein B-100 fragment.
 XX
 KW Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
 KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
 KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
 KW prothrombinase complex.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 FN W09743311-A1.
 XX
 XX 20-NOV-1997.
 XX
 PF 09-MAY-1997; 97WO-GB001255.
 XX
 PR 09-MAY-1996; 96GB-00009702.
 XX
 XX (UNLO) ROYAL FREE HOSPITAL SCHOOL MED.
 PA Bruckdorfer KR, Ettelaie C;
 PI
 XX WPI; 1998-008798/01.
 DR
 XX Peptide fragments of apo:apo:protein B-100 with anticoagulant activity -

PT used for treating or preventing coagulation, inhibiting angiogenesis,
 PT cell differentiation and apoptosis.
 XX Disclosure; Page 22; 60pp; English.
 XX
 CC This sequence is an example of the peptide of the invention. It has the
 CC formula (I), or their variants with one or more internal deletions,
 CC insertions or substitutions, while retaining anti-coagulant properties of
 CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KKQKRRH-X2-T-22 (I) X1 = S or
 CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
 CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
 CC aa. Compositions containing the peptide are used for simultaneous,
 CC separate or sequential treatment of cancer, particularly to prevent
 CC metastatic spread. They are also used to inhibit thromboplastin-mediated
 CC processes, specifically to prevent or reduce blood coagulation (e.g.
 CC during or after surgery or in cases of heart attack, stroke etc.) and to
 CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
 CC which is active as such or as part of a 98-aa peptide, inhibits
 CC activation of the prothrombinase complex; and prevents activation of
 CC factor VII on the surface of thromboplastin and of platelets by thrombin.
 CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
 CC smaller than apoB-100, they act more quickly
 XX
 SQ Sequence 15 AA;
 Query Match 71.3%; Score 38.5; DB 2; Length 15;
 Best Local Similarity 90.9%; Pred. No. 4.2;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRKRGK 11
 Db 1 TRLTRK-RGLK 10
 RESULT 10
 AAW96892
 ID AAW96892 standard; peptide; 15 AA.
 XX
 AC AAW96892;
 XX
 DT 22-APR-1999 (first entry)
 XX
 DE ApoB-100 nuclear localisation signal sequence, residues 3353-3367.
 XX
 KW Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
 XX
 OS Homo sapiens.
 XX
 FN W09856938-A1.
 XX
 XX 17-DEC-1998.
 XX
 PF 10-JUN-1998; 98WO-US011927.
 XX
 PR 13-JUN-1997; 97US-00874807.
 PR 14-MAY-1998; 98US-00079030.
 XX
 XX (BAYU) BAYLOR COLLEGE MEDICINE.
 XX
 XX Guevara JG, Hoogveen RC, Moore JP;
 XX WPI; 1998-070331/06.
 DR
 XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.
 XX
 XX Claim 19; Fig 13D; 293pp; English.
 PS
 XX AAW96878-97 represent nuclear localisation signal sequence derived from

CC human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein
 CC component of very-low density lipoproteins (VLDL), intermediate density
 CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The
 CC present sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis
 XX
 SQ Sequence 15 AA;

Query Match 71.3%; Score 38.5; DB 2; Length 15;

Best Local Similarity 90.9%; Pred. No. 4.2; Mismatches 0; Indels 1; Gaps 1;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKRGGLK 11
 DB 6 TRLTRK-RGLK 15
 ||||| |||||

RESULT 11
 ABJ37575
 ID ABJ37575 standard; peptide; 20 AA.

AC ABJ37575;

DT 10-MAY-2003 (first entry)

DE Heparin binding peptide sequence #28.

CC Cystatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;
 CC cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;
 CC rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.

OS Unidentified.

FN WO2003007689-A2.

PD 30-JAN-2003.

PF 22-JUL-2002; 2002WO-US023419.

PR 20-JUL-2001; 2001US-0306726P.

PA (ETHZ-) ETH ZUERICH.

PA (UYZU-) UNIV ZURICH.

PI Hubbell JA, Schoenmakers R, Maynard HD;

DR WPI; 2003-300420/29.

CC Use of a ligand comprising of at least one sulfated or sulfonated amino
 CC acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic
 CC retinopathy and hypoxia.

PS Disclosure; Fig 2; 79pp; English.

CC The invention relates to a novel ligand for binding a target biomolecule,
 CC which comprises a peptide having at least one sulphated or sulphonated
 CC amino acid and at least one amino acid chosen from neutral and positively
 CC charged amino acids. The novel ligands can be used for the treatment of
 CC e.g. tumours, rheumatoid arthritis, diabetic retinopathy and hypoxia.
 CC This sequence represents a heparin binding peptide relating to the
 CC invention

SQ Sequence 20 AA;

Query Match 71.3%; Score 38.5; DB 6; Length 20;

Best Local Similarity 90.9%; Pred. No. 5.5; Mismatches 0; Indels 1; Gaps 1;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKRGGLK 11
 DB 6 TRLTRK-RGLK 15
 ||||| |||||

QY 1 TRLTRKRGGLK 11
 DB 7 TRLTRK-RGLK 16
 ||||| |||||

RESULT 12

AAW57208
 ID AAW57208 standard; peptide; 22 AA.

AC AAW57208;

DT 03-AUG-1998 (first entry)

DE Apo B 100 binding site peptide analogue peptide C.

CC Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 CC growth supplement; non-natural lipid particle; low density lipoprotein;
 CC LDL; receptor component; apo B100 receptor site.

OS Synthetic.

FN Key Location/Qualifiers

FT Modified-site 1 /note= "attached to retinoic acid"

FT Modified-site 22 /note= "attached to cholesterol"

FN WO9813385-A2.

PD 02-APR-1998.

PF 25-SEP-1997; 97WO-GB002610.

PR 27-SEP-1996; 96GB-00020153.

PA (UYST) UNIV STRATHCLYDE.

PI Halbert GW, Owens MD, Baillie G;

DR WPI; 1998-230637/20.

CC Non-natural lipid particle comprising peptide binding to apo B protein
 CC receptor - useful as, e.g. vector for delivering drugs to cancer cells
 CC that express this receptor.

PS Claim 13; Fig 7; 73pp; English.

CC The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKNGRH (1) or TRLTRKGLK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor

SQ Sequence 22 AA;

Query Match 71.3%; Score 38.5; DB 2; Length 22;

Best Local Similarity 90.9%; Pred. No. 6.1; Mismatches 0; Indels 1; Gaps 1;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKRGGLK 11
 DB 7 TRLTRK-RGLK 16
 ||||| |||||

```

RESULT 13
AAW57209
ID AAW57209 standard; peptide; 22 AA.
XX
AC AAW57209;
XX
XX 03-AUG-1998 (first entry)
XX
XX Apo B 100 binding site peptide analogue peptide D.
XX
XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
KW growth supplement; non-natural lipid particle; low density lipoprotein;
KW LDL; receptor component; apo B100 receptor site.
XX
XX Synthetic.
XX
XX Key Location/Qualifiers
FT Modified-site 1
FT /note= "attached to retinoic acid"
XX
XX WO9813385-A2.
XX
XX 02-APR-1998.
XX
XX 25-SEP-1997; 97WO-GB002610.
XX
XX 27-SEP-1996; 96GB-00020153.
XX
XX (UYST ) UNIV STRATHCLYDE.
XX
XX Halbert GW, Owens MD, Baillie G;
XX
XX WPI; 1998-230637/20.
XX
XX Non-natural lipid particle comprising peptide binding to apo B protein
PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
PT that express this receptor.
XX
XX Claim 13; Fig 7; 73pp; English.
XX
XX The present sequence represents a specifically claimed Apo B 100 binding
CC site peptide analogue which can be used as a component of a non-
CC naturally occurring, receptor-competent low density lipoprotein (LDL)
CC particle of the present invention. The LDL particle comprises at least 1
CC peptide component that has at least 1 binding site for an apo B protein
CC receptor and at least 1 lipophilic substituent. Also described in the
CC invention are peptides containing an apo B binding sequence with at least
CC 70% identity with sequences: KAEYKKNKRRH (1) or TTLTRKRGK (2), or their
CC dimers. Non-naturally occurring, receptor-competent LDL particles are
CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
CC cancer cells that express an apo B protein receptor, and (ii) additives
CC for cell culture media especially as growth supplements. Non-naturally
CC occurring, receptor-competent LDL particles do not require the complete
CC apo B sequence, which is large and tends to aggregate, to provide binding
CC affinity to an apo B protein receptor
XX
XX Sequence 22 AA;
SQ
Query Match 71.3%; Score 38.5; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 6.1;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TTLTRKRGK 11
Db 7 TTLTRK-RGLK 16

RESULT 14
AAE14541
ID AAE14541 standard; peptide; 34 AA.
XX

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```

AC AAE14541;
XX
XX 17-MAY-2002 (first entry)
XX
XX Human apoB-100 derived peptide p62.
XX
XX Oxidised low density lipoprotein; oxLDL; immunoassay; periaortitis;
KW cardiovascular disease; coronary heart disease; pre-eclampsia;
KW non-insulin-dependent diabetes; endothelial dysfunction; human; apoB-100;
KW peptide p62.
XX
XX Homo sapiens.
XX
XX WO200206314-A2.
XX
XX 24-JAN-2002.
XX
XX 18-JUL-2001; 2001WO-GB003212.
XX
XX 18-JUL-2000; 2000GB-00017641.
XX
XX (ARKT-) ARK THERAPEUTICS LTD.
XX
XX Narvanen O, Yla-Herttuala S;
XX
XX WPI; 2002-179777/23.
XX
XX New peptide useful in enzyme immunoassays for detecting oxidized low
PT density lipoprotein which is a marker of coronary heart disease and other
PT cardiovascular diseases, has affinity for oxidized low density
PT lipoprotein.
XX
XX Claim 6; Page 5; 21pp; English.
XX
XX The invention relates to peptides having affinity for oxidised low
CC density lipoprotein (oxLDL), in cyclised or multimeric form. The peptide
CC is useful in an immunoassay to determine the presence, and optionally,
CC the amount of antibodies in a sample, having affinity for oxLDL.
CC Preferably immobilised peptide is useful for measuring the amount of
CC autoantibodies for oxLDL in a sample, especially a serum or plasma sample
CC from a patient for evaluating the risk of coronary heart diseases, other
CC cardiovascular diseases, and several other disorders such as
CC periaortitis, pre-eclampsia, non-insulin-dependent diabetes and
CC endothelial dysfunction. The peptide of the invention is stable, can be
CC synthesised easily without the need to isolate proteins from a patient's
CC blood, and has a long half-life. The present sequence is human apoB-100
CC derived peptide p62 used in the invention
XX
XX Sequence 34 AA;
SQ
Query Match 71.3%; Score 38.5; DB 5; Length 34;
Best Local Similarity 90.9%; Pred. No. 9.4;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TTLTRKRGK 11
Db 25 TTLTRK-RGLK 34

RESULT 15
AAW96876
ID AAW96876 standard; peptide; 36 AA.
XX
XX AAW96876;
XX
XX 22-APR-1999 (first entry)
XX
XX Nucleic acid binding domain from apoB-100, residues 3348-3390.
XX
XX Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
KW non-small cell lung carcinoma; diabetes; arteriosclerosis.

```

XX OS Homo sapiens.
 XX PN WO9856938-A1.
 XX PD 17-DEC-1998.
 XX PF 10-JUN-1998; 98WO-US011927.
 XX PR 13-JUN-1997; 97US-00874807.
 XX PR 14-MAY-1998; 98US-00079030.
 XX PA (BAYU) BAYLOR COLLEGE MEDICINE.
 XX PI Guevara JG, Hoogvee RC, Moore JP;
 XX DR WPI; 1999-070331/06.
 XX PT Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense
 PT treatment.
 XX PS Claim 16; Fig 12C; 293pp; English.
 XX CC AAW96827-77 represent nucleic acid binding domains derived from human
 CC apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein component
 CC of very-low density lipoproteins (VLDL), intermediate density lipoprotein
 CC (IDL), low density lipoproteins (LDL) and lipoprotein a. The present
 CC sequence can be used in the composition of the invention. The
 CC specification describes a composition that comprises LDL and
 CC apolipoproteins for the binding and in vivo transport of nucleic acids.
 CC The composition is used to deliver nucleic acids to eukaryotic cells, in
 CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense
 CC molecule (or ribozyme). Specifically they are used for gene therapy of
 CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic
 CC fibrosis and arteriosclerosis
 XX SQ Sequence 36 AA;
 Query Match 71.3%; Score 38.5; DB 2; Length 36;
 Best Local Similarity 90.9%; Pred. No. 9.9;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 Qy 1 TRLTRKRGGLK 11
 Db 11 TRLTRK-RGLK 20
 Search completed: January 13, 2005, 01:43:04
 Job time : 86.1011 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:30:05 ; Search time 15.8689 Seconds
(without alignments)
66.696 Million cell updates/sec

Title: US-09-823-418-17
Perfect score: 54
Sequence: 1 TRLTRKRGRLK 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79:.*
1: pir1:.*
2: pir2:.*
3: pir3:.*
4: pir4:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|----------|--------------------|
| 1 | 42 | 77.8 | 2073 | 1 BWASBE | bimE protein - Eme |
| 2 | 38.5 | 71.3 | 596 | 2 S32802 | apolipoprotein B - |
| 3 | 38.5 | 71.3 | 4563 | 1 LRHUB | apolipoprotein B-1 |
| 4 | 37 | 68.5 | 488 | 2 T49903 | glucosyltransferas |
| 5 | 36 | 66.7 | 248 | 2 H84008 | hypothetical prote |
| 6 | 36 | 66.7 | 250 | 2 D98082 | hypothetical prote |
| 7 | 36 | 66.7 | 264 | 2 A12112 | hypothetical prote |
| 8 | 36 | 66.7 | 405 | 2 C72305 | transposase, IS605 |
| 9 | 35 | 64.8 | 100 | 2 A36950 | urease (EC 3.5.1.5 |
| 10 | 35 | 64.8 | 346 | 2 S49963 | hypothetical prote |
| 11 | 35 | 64.8 | 807 | 2 AC1031 | hypothetical prote |
| 12 | 34.5 | 63.9 | 269 | 2 C60950 | apolipoprotein B-1 |
| 13 | 34.5 | 63.9 | 779 | 2 JH0102 | apolipoprotein B - |
| 14 | 34 | 63.0 | 221 | 2 B64188 | arginine transport |
| 15 | 34 | 63.0 | 250 | 2 A69843 | hypothetical prote |
| 16 | 34 | 63.0 | 608 | 2 A46312 | gag polyprotein - |
| 17 | 33 | 61.1 | 191 | 2 T25791 | hypothetical prote |
| 18 | 33 | 61.1 | 246 | 2 T38787 | hypothetical prote |
| 19 | 33 | 61.1 | 284 | 2 G82319 | DnaJ-related prote |
| 20 | 33 | 61.1 | 305 | 1 NKVLHH | core antigen - her |
| 21 | 33 | 61.1 | 332 | 2 T13447 | hypothetical prote |
| 22 | 33 | 61.1 | 541 | 2 AF0547 | propionate catabol |
| 23 | 33 | 61.1 | 605 | 2 T04197 | hypothetical prote |
| 24 | 33 | 61.1 | 643 | 2 AF2471 | penicillin-binding |
| 25 | 33 | 61.1 | 710 | 2 T46589 | copy-2 protein (im |
| 26 | 32.5 | 60.2 | 275 | 2 E60950 | apolipoprotein B-1 |
| 27 | 32 | 59.3 | 100 | 2 E7102 | urease (EC 3.5.1.5 |
| 28 | 32 | 59.3 | 132 | 2 G75409 | hypothetical prote |
| 29 | 32 | 59.3 | 150 | 2 H72645 | hypothetical prote |

| | | | | | | |
|----|----|------|-----|---|---------|--------------------|
| 30 | 32 | 59.3 | 202 | 2 | G84502 | hypothetical prote |
| 31 | 32 | 59.3 | 220 | 2 | A75287 | response regulator |
| 32 | 32 | 59.3 | 250 | 2 | F95218 | hypothetical prote |
| 33 | 32 | 59.3 | 266 | 2 | AB1827 | hypothetical prote |
| 34 | 32 | 59.3 | 301 | 2 | AF2223 | heterodisulfide re |
| 35 | 32 | 59.3 | 345 | 2 | T48758 | hypothetical prote |
| 36 | 32 | 59.3 | 398 | 2 | S40752 | hypothetical prote |
| 37 | 32 | 59.3 | 429 | 2 | E83723 | hypothetical prote |
| 38 | 32 | 59.3 | 441 | 2 | JQ2191 | nucleosid prote |
| 39 | 32 | 59.3 | 458 | 2 | D70410 | cytosolic axial fi |
| 40 | 32 | 59.3 | 627 | 2 | T00484 | hypothetical prote |
| 41 | 32 | 59.3 | 637 | 2 | JH0611 | glutamate-cysteine |
| 42 | 32 | 59.3 | 637 | 2 | AJ35015 | hypothetical prote |
| 43 | 32 | 59.3 | 771 | 2 | T29177 | hypothetical prote |
| 44 | 32 | 59.3 | 798 | 2 | AI2053 | competence protein |
| 45 | 32 | 59.3 | 844 | 2 | S05988 | translation elonga |

ALIGNMENTS

RESULT 1

BWASBE

bimE protein - Emericella nidulans

C:Species: Emericella nidulans, Aspergillus nidulans

C:Date: 31-Dec-1991 #sequence_revision 31-Dec-1991 #text_change 09-Jul-2004

C:Accession: A37879

R:Engle, D.B.; Osmani, S.A.; Osmani, A.H.; Rosborough, S.; Xiang, X.; Morris, N.R.

J. Biol. Chem. 265, 16132-16137, 1990

A:Title: A negative regulator of mitosis in Aspergillus is a putative membrane-spanning

A:Reference number: A37879; MUID:90375468; PMID:1697851

A:Accession: A37879

A:Molecule type: mRNA

A:Residues: 1-2073 <ENG>

A:Cross-references: UNIPROT:P24686; GB:M59705; GB:J05607; NID:g168026; PIDN:AAA51478.1;

A>Note: In addition to three predicted transmembrane domains, there are several potent

asein kinase, and one sequence that resembles a nuclear localization signal

C:Comment: This protein is part of a regulatory pathway that includes the nimA protein

ter mitosis and prevent them from leaving mitosis.

C:Genetics:

A:Gene: bimE

C:Superfamily: bimE protein

C:Keywords: cell cycle control; mitosis; transmembrane protein

F:1623-1643/Domain: transmembrane #status predicted <TM1>

F:1685-1703/Domain: transmembrane #status predicted <TM2>

F:1746-1764/Domain: transmembrane #status predicted <TM3>

Query Match 77.8%; Score 42; DB 1; Length 2073;

Best Local Similarity 80.0%; Pred. No. 14;

Matches 8; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TRLTRKRGRL 10

Db 832 TRLTRKRGRL 841

RESULT 2

S32802

apolipoprotein B - crab-eating macaque (fragment)

C:Species: Macaca fascicularis (crab-eating macaque)

C:Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Jul-2004

C:Accession: S32802

R:Pape, M.E.; Castle, C.K.; Murray, R.W.; Funk, G.M.; Hunt, C.E.; Marotti, K.R.; Melch

Biochim. Biophys. Acta 1086, 326-334, 1991

A:Title: Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional

A:Reference number: S32802; MUID:92075708; PMID:1742325

A:Accession: S32802

A>Status: preliminary

A:Molecule type: mRNA

A:Residues: 1-596 <PAP>

A:Cross-references: UNIPROT:Q28473; EMBL:X15737; NID:g38047; PIDN:CAA33755.1; PID:g9301

C:Superfamily: apolipoprotein B

| | | | | | |
|---|--------|------------------|-------|-------------|---------|
| Query Match | 71.3%; | Score 38.5; | DB 2; | Length 596; | |
| Best Local Similarity | 90.9%; | Pred. No. 21; | | | |
| Matches | 10; | Conservative | 0; | Mismatches | 1; Gaps |
| | | | | | |
| Qv | 1 | TLRLTRKRGGLK 11 | | | |
| | | | | | |
| Db | 226 | TLRLTRK-RGLK 235 | | | |
| RESULT 3 | | | | | |
| LPHUB | | | | | |
| apolipoprotein B-100 precursor - human | | | | | |
| N:Contains: apolipoprotein B-26; apolipoprotein B-48; apolipoprotein B-74 | | | | | |
| C:Species: Homo sapiens (man) | | | | | |
| C:Date: 28-Dec-1987 #sequence revision 28-Dec-1987 #text change 09-Jul-2004 | | | | | |
| C:Accession: A27850; A25679; A25263; A25267; A24320; A24684; A23817; A25774; A28452; I61909; I59510; I39474; I39469; I84624; I37179; PS0058 | | | | | |
| R:Ludwig, E.H.; Blackhart, B.D.; Pierotti, V.R.; Caiati, L.; Fortier, C.; Knott, T.; Soc | | | | | |
| DNA 6, 363-372, 1987 | | | | | |
| A:Title: DNA sequence of the human apolipoprotein B gene. | | | | | |
| A:Reference number: A27850; MUID:86003974; PMID:3652907 | | | | | |
| A:Accession: A27850 | | | | | |
| A:Molecule type: DNA | | | | | |
| A:Residues: 1-617, 'A', 619-1929, 'F', 1931-3318, 'D', 3320-3426, 'T', 3428-3431, 'Q', 3433-3731, 'A' | | | | | |
| A:Cross-references: UNIPROT:P04114; UNIPROT:P78482; UNIPROT:P78479; UNIPROT:Q9UMN0; UNIB | | | | | |
| R:Cladaras, C.; Hadzopoulou-Cladaras, M.; Nolte, R.T.; Atkinson, D.; Zannis, V.I. | | | | | |
| EMBO J. 5, 3495-3507, 1986 | | | | | |
| A:Title: The complete sequence and structural analysis of human apolipoprotein B-100: re | | | | | |
| A:Reference number: A91058; MUID:87161758; PMID:3030729 | | | | | |
| A:Accession: A25679 | | | | | |
| A:Molecule type: mRNA | | | | | |
| A:Residues: 1-11, 15-2539, 'S', 2541-3823, 'R', 3825-4563 <CLA> | | | | | |
| A>Note: 1109-Aap was also found | | | | | |
| R:Knott, T.J.; Wallis, S.C.; Powell, L.M.; Pease, R.J.; Lusis, A.J.; Blackhart, B.; McCa | | | | | |
| Nucleic Acids Res. 14, 7501-7503, 1986 | | | | | |
| A:Title: Complete cDNA and derived protein sequence of human apolipoprotein B-100. | | | | | |
| A:Reference number: A93639; MUID:87016385; PMID:3763409 | | | | | |
| A:Accession: A25263 | | | | | |
| A:Molecule type: mRNA | | | | | |
| A:Residues: 1-272, 'N', 274-617, 'A', 619-1217, 'E', 1219-2091, 'V', 2093-2364, 'T', 2366-2679, 'Q' | | | | | |
| A:Cross-references: GB:M04506; NID:G34330; PIDN:CAA28191.1; PID:G34331 | | | | | |
| R:Law, S.W.; Grant, S.M.; Higuchi, K.; Hospattankar, A.; Lackner, K.; Lee, N.; Brewer JH | | | | | |
| Proc. Natl. Acad. Sci. U.S.A. 83, 8142-8146, 1986 | | | | | |
| A:Title: Human liver apolipoprotein B-100 cDNA: complete nucleic acid and derived amino | | | | | |
| A:Reference number: A94134; MUID:87041416; PMID:3464946 | | | | | |
| A:Accession: A25267 | | | | | |
| A:Molecule type: mRNA | | | | | |
| A:Residues: 1-617, 'A', 619-703, 'P', 705-792, 'R', 794-1270, 'S', 1272-1866, 'G', 1868-2036, 'N', 2 | | | | | |
| 4189-4220, 'M', 4222-4563 <LAW> | | | | | |
| A>Note: the codons given for residues 704, 793, 1271, 2037, 2933, 3286, 3782, 4188, and | | | | | |
| R:Chen, S.H.; Yang, C.Y.; Chen, P.P.; Setzer, D.; Tanimura, M.; Li, W.H.; Gotto Jr., A.M | | | | | |
| J. Biol. Chem. 261, 12918-12921, 1986 | | | | | |
| A:Title: The complete cDNA and amino acid sequence of human apolipoprotein B-100. | | | | | |
| A:Reference number: A92556; MUID:87008488; PMID:3759943 | | | | | |
| A:Accession: A25266 | | | | | |
| A:Molecule type: mRNA | | | | | |
| A:Residues: 1-97, 'I', 99-328, 'V', 330-644, 'I', 646-918, 'P', 920-3318, 'D', 3320-3426, 'T', 3428- | | | | | |
| 9-4132, 'G', 4134-4180, 'E', 4182-4563 <CHE> | | | | | |
| A:Cross-references: GB:J02610; NID:G178803; PIDN:AAA35549.1; PID:G178804 | | | | | |
| A>Note: a total of 2366 residues were confirmed by direct sequencing of tryptic peptides | | | | | |
| R:Protter, A.A.; Hardman, D.A.; Sato, K.Y.; Schilling, J.W.; Yamanaka, M.; Hort, Y.J.; H | | | | | |
| Proc. Natl. Acad. Sci. U.S.A. 83, 5678-5682, 1986 | | | | | |
| A:Title: Analysis of cDNA clones encoding the entire B-26 region of human apolipoprotein | | | | | |
| A:Reference number: A24320; MUID:86287319; PMID:3461454 | | | | | |
| A:Accession: A24320 | | | | | |
| A:Molecule type: mRNA | | | | | |
| A:Residues: 1-97, 'I', 99-617, 'A', 619-941, 'Y'YTWSLPKP', 951-1138, 'PTGRLPNCFNGSLICYSLWLHSPQE | | | | | |
| A:Cross-references: GB:M14081; NID:G178795; PIDN:AAA51752.1; PID:G553189 | | | | | |
| R:Law, S.W.; Lackner, K.J.; Hospattankar, A.V.; Anchors, J.M.; Sakaguchi, A.Y.; Naylor, | | | | | |
| Proc. Natl. Acad. Sci. U.S.A. 82, 8340-8344, 1985 | | | | | |
| A:Title: Human apolipoprotein B-100: cloning, analysis of liver mRNA, and assignment of | | | | | |
| A:Reference number: A24684; MUID:86094221; PMID:3001697 | | | | | |
| A:Accession: A24684 | | | | | |

A:Molecule type: mRNA
A:Residues: 485-617, 'A', 619-1044 <LA2>
A:Cross-references: GB:M12480; NID:G178791; PIDN:AAA51751.1; PID:G178792
R:Protter, A.A.; Hardman, D.A.; Schilling, J.W.; Miller, J.; Appleby, V.; Chen, G.C.; K
Proc. Natl. Acad. Sci. U.S.A. 83, 1467-1471, 1986
A:Title: Isolation of a cDNA clone encoding the amino-terminal region of human apolipop
A:Reference number: A94088; MUID:86149325; PMID:3513177
A:Accession: A23817
A:Molecule type: mRNA
A:Residues: 1-291 <PRO>
A:Cross-references: GB:M12681; NID:G178797; PIDN:AAA51753.1; PID:G178798
R:Deeb, S.S.; Motulsky, A.G.; Albers, J.J.
Proc. Natl. Acad. Sci. U.S.A. 82, 4983-4986, 1985
A:Title: A partial cDNA clone for human apolipoprotein B.
A:Reference number: A25774; MUID:85270450; PMID:3860836
A:Accession: A25774
A:Molecule type: mRNA
A:Residues: 709-791, 'SSSWKAASHGCPHSAGD', 810-906 <DEE>
A:Cross-references: GB:K03175; NID:G178821; PIDN:AAA51759.1; PID:G178822
R:Carlsson, P.; Darnfors, C.; Olofsson, S.O.; Bjursell, G.
Gene 49, 29-51, 1986
A:Title: Analysis of the human apolipoprotein B gene; complete structure of the B-74 re
A:Reference number: A91565; MUID:87191999; PMID:2883086
A:Accession: A26533
A:Molecule type: mRNA
A:Residues: 1282-2721, 2742-3290, 'L', 3292-3336, 'N', 3338-3948, 'F', 3950-3963, 'Y', 3965-4180
A:Cross-references: GB:M15421; NID:G178817; PIDN:AAA51758.1; PID:G178818
R:Hardman, D.A.; Protter, A.A.; Chen, G.C.; Schilling, J.W.; Sato, K.Y.; Lau, K.; Yaman
Biochemistry 26, 5478-5486, 1987
A:Title: Structural comparison of human apolipoproteins B-48 and B-100.
A:Reference number: A29671; MUID:88050832; PMID:3676265
A:Accession: A29671
A:Molecule type: mRNA
A:Residues: 1671-2323, 'PYW', 2327-2352, 'H', 2354-2398 <HAR>
A:Cross-references: GB:M17367; NID:G178731; PIDN:AAA51741.1; PID:G178732
R:Shoulders, C.C.; Myant, N.B.; Sidoli, A.; Rodriguez, J.C.; Cortese, C.; Baralle, F.E
Atherosclerosis 58, 277-289, 1985
A:Title: Molecular cloning of human LDL apolipoprotein B cDNA. Evidence for more than o
A:Reference number: A90084; MUID:86130855; PMID:3841481
A:Accession: A29287
A:Molecule type: mRNA
A:Residues: 3846-4298 <SHO>
R:Pfizner, R.; Wagener, R.; Stoffel, W.
Biol. Chem. Hoppe-Seyler 367, 1077-1083, 1986
A:Title: Isolation, expression and characterization of a human apolipoprotein B 100-spe
A:Reference number: A25572; MUID:87076044; PMID:3024665
A:Accession: A25572
A:Molecule type: mRNA
A:Residues: 4219-4337, 'S', 4339-4563 <PFI>
A:Cross-references: GB:M36676
R:Wei, C.F.; Chen, S.H.; Yang, C.Y.; Marcel, Y.L.; Milne, R.W.; Li, W.H.; Sparrow, J.T.
Proc. Natl. Acad. Sci. U.S.A. 82, 7265-7269, 1985
A:Reference number: A24738; MUID:86042646; PMID:2932736
A:Accession: A24738
A:Molecule type: mRNA
A:Residues: 'N', 3729-3731, 'I', 3733-3875, 'A', 3877-3948, 'F', 3950-3963, 'Y', 3965-3982, 'S', 3
A:Cross-references: GB:M12413; NID:G178735; PIDN:AAA51742.1; PID:G178736
R:Chen, S.H.; Habib, G.; Yang, C.Y.; Gu, Z.W.; Lee, B.R.; Weng, S.; Silberman, S.R.; Ca
Science 238, 363-366, 1987
A:Title: Apolipoprotein B-48 is the product of a messenger RNA with an organ-specific i
A:Reference number: A40133; MUID:88018019; PMID:3659919
A:Accession: B40133
A:Molecule type: mRNA
A:Residues: 2165-2179 <CHI>
A:Cross-references: GB:M18036; NID:G178799; PIDN:AAA51754.1; PID:G178800
A>Note: this mRNA includes the stop codon of the organ-specific mRNA for apo48
A:Accession: A40133
A:Molecule type: protein
A:Residues: 51-75; 101-110, 129-139; 158-174, 197-207; 276-287; 298-304; 306-314; 526-532; 538-5
36; 1486-1498; 1537-1556; 1563-1572; 1601-1610; 1647-1661; 1697-1724; 1770-1781; 1859-1897; 1968
A>Note: these fragments were derived from apo48
R:Hardman, D.A.; Protter, A.A.; Schilling, J.W.; Kane, J.P.
Biochem. Biophys. Res. Commun. 149, 1214-1219, 1987

A;Title: Carboxyl terminal analysis of human B-48 protein confirms the novel mechanism of
A;Reference number: A28002; MUID:88106542; PMID:3426612
A;Accession: A28002
A;Molecule type: mRNA
A;Residues: 2129-2179, 2181-2235 <HA2>
A;Cross-references: GB:M18471
A;Experimental source: intestine
A;Note: this mRNA from intestine includes a stop codon created by RNA editing in place of
R;Mehrabian, M.; Schumaker, V.N.; Fareed, G.C.; West, R.; Johnson, D.F.; Kirchgessner, T.
Nucleic Acids Res. 13, 6937-6953, 1985
A;Title: Human apolipoprotein B: identification of cDNA clones and characterization of m
A;Reference number: A24269; MUID:86041888; PMID:3903660
A;Accession: A24269
A;Molecule type: mRNA
A;Residues: 3056-3159 <MEH>
A;Cross-references: GB:X03045; NID:928783; PID:CAA26850.1; PID:9929609
R;Hospatankar, A.V.; Higuchi, K.; Law, S.W.; Meglin, N.; Brewer Jr., H.B.
Biochem. Biophys. Res. Commun. 148, 279-285, 1987
A;Title: Identification of a novel in-frame translational stop codon in human intestine
A;Reference number: A29659; MUID:88049670; PMID:2445342
A;Accession: A29659
A;Molecule type: mRNA
A;Residues: 2169-2179 <HOS>
A;Note: the sequence shown represents the carboxyl end of apolipoprotein B-48
A;Note: two RNA species, 14.1kb and 7.5kb in length, were isolated from the human intest
ch encodes the 250K apob-48, CAA encoding 2180-Gln is substituted by the stop codon TAA,
R;Yang, C.; Kim, T.W.; Weng, S.; Lee, B.; Yang, M.; Gotto Jr., A.M.
Proc. Natl. Acad. Sci. U.S.A. 87, 5523-5527, 1990
A;Title: Isolation and characterization of sulfhydryl and disulfide peptides of human ap
A;Reference number: A35783; MUID:90319144; PMID:2115173
A;Contents: disulfide bonds
A;Accession: A35783
A;Molecule type: protein
A;Residues: 28-41;76-97, 'I', 99-100;175-193;206-215;239-249;259-266;357-399;455-490;512-5
A;Note: cysteines at positions 112, 1422, 1505, 1662, 3761, 3917, and 4217 have free su
R;LeBoeuf, R.C.; Miller, C.; Shively, J.E.; Schumaker, V.N.; Balla, M.A.; Lusis, A.J.
FEBS Lett. 170, 105-108, 1984
A;Title: Human apolipoprotein B: partial amino acid sequence.
A;Reference number: A22006; MUID:84208786; PMID:6373369
A;Accession: A22006
A;Molecule type: protein
A;Residues: 873-892, 'K', 894-896 <LE1>
A;Accession: B22006
A;Molecule type: protein
A;Residues: 3113, 'L', 3115-3130, 'R', 3132-3133, 'P', 3135-3136, 'R' <LE2>
R;Blackhart, B.D.; Ludwig, E.M.; Pierotti, V.R.; Caiati, L.; Onasch, M.A.; Wallis, S.C.;
J. Biol. Chem. 261, 15364-15367, 1986
A;Title: Structure of the human apolipoprotein B gene.
A;Reference number: A92564; MUID:87057153; PMID:2946672
A;Contents: annotation; gene structure
R;Wagener, R.; Pfitzner, R.; Stoffel, W.
Biol. Chem. Hoppe-Seyler 369, 419-425, 1987
A;Title: Studies on the organization of the human apolipoprotein B 100 gene.
A;Reference number: A90715; MUID:87271140; PMID:2886136
A;Contents: annotation; gene structure
R;Weisgraber, K.H.; Rall Jr., S.C.
J. Biol. Chem. 262, 11097-11103, 1987
A;Title: Human apolipoprotein B-100 heparin-binding sites.
A;Reference number: A92605; MUID:87280197; PMID:3301850
A;Contents: annotation; heparin binding and disulfide bond
R;Dashit, N.; Lee, D.M.; Mok, T.
Biochem. Biophys. Res. Commun. 137, 493-499, 1986
A;Title: Apolipoprotein B is a calcium binding protein.
A;Reference number: A90125; MUID:86242245; PMID:3087360
A;Contents: annotation; calcium binding
R;Carlsson, P.; Olofsson, S.O.; Bondjers, G.; Darnfors, C.; Wiklund, O.; Bjursell, G.
Nucleic Acids Res. 13, 8813-8826, 1985
A;Title: Molecular cloning of human apolipoprotein B cDNA.
A;Reference number: I37178; MUID:86093680; PMID:3841204
A;Accession: I37180

Query Match

71.3%; Score 38.5; DB 1; Length 4563;

Best Local Similarity 90.9%; Pred. No. 1.3e+02;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY 1 LTRTKRERGLK 11
DB 3385 LTRTKR-RGLK 3394
RESULT 4
T49903
glucosyltransferase-like protein - Arabidopsis thaliana
N;Alternate names: protein T24H18.60
C;Species: Arabidopsis thaliana (mouse-ear cress)
C;Date: 02-Jun-2000 #sequence_revision 02-Jun-2000 #text_change 09-Jul-2004
C;Accession: T49903
R;Bavan, M.; Robben, J.; Grymonprez, B.; Volckaert, G.; Bancroft, I.; Mewes, H.W.; Rudi
submitted to the Protein Sequence Database, April 2000
A;Reference number: Z25024
A;Accession: T49903
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-488 <BEV>
A;Cross-references: UNIPROT:Q9LXV0; EMBL:AL353013; GSPDB:GN00063; ATSP:T24H18.60
C;Experimental source: cultivar Columbia; BAC clone T24H18
C;Genetics:
A;Gene: ATSP:T24H18.60
A;Map position: 5
C;Superfamily: flavonol O3-glucosyltransferase
Query Match 68.5%; Score 37; DB 2; Length 488;
Best Local Similarity 77.8%; Pred. No. 34;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;
QY 2 RLTRKERGL 10
DB 344 RLTRSERGL 352
RESULT 5
H84008
hypothetical protein BH2872 [imported] - Bacillus halodurans (strain C-125)
C;Species: Bacillus halodurans
C;Date: 01-Dec-2000 #sequence_revision 01-Dec-2000 #text_change 09-Jul-2004
C;Accession: H84008
R;Takami, H.; Nakasone, K.; Takaki, Y.; Maeno, G.; Sasaki, R.; Masui, N.; Fujii, F.; Hir
Nucleic Acids Res. 28, 4317-4331, 2000
A;Title: Complete genome sequence of the alkaliphilic bacterium Bacillus halodurans and
A;Reference number: A83650; MUID:20512582; PMID:11058132
A;Accession: H84008
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-248 <STO>
A;Cross-references: UNIPROT:Q9K8Y0; GB:AP001516; GB:BA000004; NID:G10175192; PIDN:BA506
A;Experimental source: strain C-125
C;Genetics:
A;Gene: BH2872
C;Superfamily: Bacillus subtilis hypothetical protein yjba
Query Match 66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 28;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 3 LTRKERGLK 11
DB 155 LTRKERQLK 163
RESULT 6
D98082
hypothetical protein fecE [imported] - Streptococcus pneumoniae (strain R6)
C;Species: Streptococcus pneumoniae
C;Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 09-Jul-2004
C;Accession: D98082

A;Experimental source: strain MSB8
C;Genetics:
A;Gene: TM1044
C;Superfamily: hypothetical protein b1432

Query Match 66.7%; Score 36; DB 2; Length 405;
Best Local Similarity 70.0%; Pred. No. 44;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLTRKERGLK 11
||:|:|:|
Db 227 RLSRKQSGK 236

RESULT 9
A36950
urease (EC 3.5.1.5) 11k chain - Bacillus sp. (strain TB-90)
N;Alternate names: urea protein
C;Species: Bacillus sp.
C;Date: 07-Jul-1995 #sequence_revision 07-Jul-1995 #text_change 09-Jul-2004
C;Accession: A36950
R;Maeda, M.; Hidaka, M.; Nakamura, A.; Maeaki, H.; Uozumi, T.
J. Bacteriol. 176, 432-442, 1994
A;Title: Cloning, sequencing, and expression of thermophilic Bacillus sp. strain TB-90
A;Reference number: A36950; MUID:94117379; PMID:8288539
A;Accession: A36950
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-100 <MAP>
A;Cross-references: UNIPROT:Q07399; GB:D14439; NID:G393296; PIDN:BAA03323.1; PID:G2163
C;Superfamily: urease, gamma subunit; urease 11k chain homology
C;Keywords: hydrolase
F;1-100/Domain: urease 11k chain homology <U11>

Query Match 64.8%; Score 35; DB 2; Length 100;
Best Local Similarity 100.0%; Pred. No. 19;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 RKERGLK 11
|||:|:|
Db 23 RKERGLK 29

RESULT 10
S49963
hypothetical protein Y1L019w - yeast (Saccharomyces cerevisiae)
N;Alternate names: hypothetical protein Y13299.12
C;Species: Saccharomyces cerevisiae
C;Date: 28-May-1993 #sequence_revision 24-Feb-1995 #text_change 09-Jul-2004
C;Accession: S49963
R;Skellton, J. J. Churcher, C.
submitted to the EMBL Data Library, December 1994
A;Reference number: S49951
A;Accession: S49963
A;Molecule type: DNA
A;Residues: 1-146 <SKE>
A;Cross-references: UNIPROT:P40546; EMBL:Z46881; NID:G599967; PIDN:CAA86973.1; PID:G599967
C;Genetics:
A;Gene: MIPS:Y1L019w
A;Cross-references: SGD:S0001281
A;Map position: 9L

Query Match 64.8%; Score 35; DB 2; Length 346;
Best Local Similarity 60.0%; Pred. No. 59;
Matches 6; Conservative 3; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLTRKERGLK 11
:|:|:|:|
Db 298 KATRKERGLK 307

RESULT 11
AC1031

A;Authors: Schleich, S.; Schroeter, R.; Scoffone, F.; Sekiguchi, J.; Sekowska, A.; Serod
akeuchi, M.; Tanakoshi, A.; Tanaka, T.; Terpstra, P.; Tognoni, A.; Tosato, V.; Uchiyama,
T.; Winters, P.; Wipat, A.; Yamamoto, H.; Yamane, K.; Yasunoto, K.; Yata, K.; Yoshida, K.
A;Authors: Yoshikawa, H.F.; Zumstein, E.; Yoshikawa, H.; Danchin, A.
A;Title: The complete genome sequence of the Gram-positive bacterium *Bacillus subtilis*.
A;Reference number: A69580; MUID:98044033; PMID:9384377
A;Accession: A69843
A;Status: preliminary; nucleic acid sequence not shown; translation not shown
A;Molecule type: DNA
A;Residues: 1-250 <KUN>
A;Cross-references: UNIPROT:O31597; GB:Z99110; GB:AL009126; NID:g2633472; PIDN:CAB12998.
A;Experimental source: strain 168
C;Genetics:
A;Gene: yjba
C;Superfamily: *Bacillus subtilis* hypothetical protein yjba

Query Match 63.0%; Score 34; DB 2; Length 250;
Best Local Similarity 77.8%; Pred. NO. 69;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LTRKERGLK 11
| | | | | : |
Db 157 LTRKERQIK 165

Search completed: January 13, 2005, 01:52:42
Job time : 17.8689 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 85.8361 seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-17
Perfect score: 54
Sequence: 1 TRLTRKRGK 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : UniProt_02.*
1: uniprot_sprot.*
2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|--------------|--------------------|
| 1 | 42 | 77.8 | 2073 | 1 BIME_EMENI | P24686 emericella |
| 2 | 38.5 | 71.3 | 414 | 2 Q7YQR5 | Q7YQR5 actus vocif |
| 3 | 38.5 | 71.3 | 596 | 2 Q28473 | Q28473 macaca fasc |
| 4 | 38.5 | 71.3 | 3262 | 2 Q13788 | Q13788 homo sapien |
| 5 | 38.5 | 71.3 | 4563 | 1 ABE_HUMAN | P04114 homo sapien |
| 6 | 38.5 | 71.3 | 4563 | 2 Q7Z600 | Q7Z600 homo sapien |
| 7 | 37 | 68.5 | 431 | 2 Q7U8V5 | Q7U8V5 synecococc |
| 8 | 37 | 68.5 | 488 | 2 Q9LXV0 | Q9LXV0 arabidopsis |
| 9 | 37 | 68.5 | 488 | 2 Q8VZF9 | Q8VZF9 arabidopsis |
| 10 | 36 | 66.7 | 94 | 2 Q89I78 | Q89I78 bradyrhizob |
| 11 | 36 | 66.7 | 248 | 2 Q8HM04 | Q8HM04 bacillus th |
| 12 | 36 | 66.7 | 248 | 2 Q73BX2 | Q73BX2 bacillus ce |
| 13 | 36 | 66.7 | 248 | 2 Q81GL8 | Q81GL8 bacillus ce |
| 14 | 36 | 66.7 | 248 | 2 Q81TS7 | Q81TS7 bacillus an |
| 15 | 36 | 66.7 | 248 | 2 Q9K8Y0 | Q9K8Y0 bacillus ha |
| 16 | 36 | 66.7 | 248 | 2 AAS40225 | AAS40225 bacillus |
| 17 | 36 | 66.7 | 248 | 2 AAT30275 | AAT30275 bacillus |
| 18 | 36 | 66.7 | 250 | 2 Q8DNJ3 | Q8DNJ3 streptococc |
| 19 | 36 | 66.7 | 252 | 2 Q8ERU3 | Q8ERU3 oceanobacil |
| 20 | 36 | 66.7 | 264 | 2 Q8YU98 | Q8YU98 anabaena sp |
| 21 | 36 | 66.7 | 374 | 2 Q8YU98 | Q8YU98 anabaena sp |
| 22 | 36 | 66.7 | 378 | 2 Q7P868 | Q7P868 fusobacteri |
| 23 | 36 | 66.7 | 378 | 2 Q8RIN6 | Q8RIN6 fusobacteri |
| 24 | 36 | 66.7 | 405 | 2 Q9X0D4 | Q9X0D4 thermotoga |
| 25 | 36 | 66.7 | 1095 | 1 A79B_MOUSE | P98195 mus musculu |
| 26 | 35 | 64.8 | 79 | 2 Q6ZE28 | Q6ZE28 oryza sativ |
| 27 | 35 | 64.8 | 79 | 2 BAC83466 | BAC83466 oryza sat |
| 28 | 35 | 64.8 | 100 | 1 URE3_BACSB | Q73399 bacillus sp |
| 29 | 35 | 64.8 | 100 | 2 Q733J4 | Q733J4 bacillus ce |
| 30 | 35 | 64.8 | 100 | 2 AAS42569 | AAS42569 bacillus |
| 31 | 35 | 64.8 | 117 | 2 Q82B17 | Q82B17 streptomyce |

| | | | | | |
|----|----|------|-----|--------------|---------------------|
| 32 | 35 | 64.8 | 210 | 2 Q7RYL3 | Q7ryl3 neurospora |
| 33 | 35 | 64.8 | 259 | 2 Q88JG3 | Q88jg3 pseudomonas |
| 34 | 35 | 64.8 | 281 | 2 Q72VN4 | Q72vn4 leptospira |
| 35 | 35 | 64.8 | 281 | 2 Q8F989 | Q8f989 leptospira |
| 36 | 35 | 64.8 | 281 | 2 AAS68890 | AAS68890 leptospir |
| 37 | 35 | 64.8 | 322 | 2 Q6CU76 | Q6cu76 kluyveromyc |
| 38 | 35 | 64.8 | 340 | 2 Q8GLA1 | Q8glal streptococc |
| 39 | 35 | 64.8 | 345 | 1 DDL_WOLSU | Q7ma71 wolinnella s |
| 40 | 35 | 64.8 | 346 | 1 YIB9_YEAST | P40546 saccharomyc |
| 41 | 35 | 64.8 | 346 | 2 Q6Q5C5 | Q6q5g5 saccharomyc |
| 42 | 35 | 64.8 | 346 | 2 AAS56415 | AAS56415 saccharom |
| 43 | 35 | 64.8 | 401 | 2 Q7S6B9 | Q7s6b9 neurospora |
| 44 | 35 | 64.8 | 438 | 2 Q99JK2 | Q99jk2 mus musculu |
| 45 | 35 | 64.8 | 439 | 2 Q8RUL7 | Q8rul7 oryza sativ |

ALIGNMENTS

RESULT 1
BIME EMENI STANDARD; PRT; 2073 AA.
AC P24686;
DT 01-MAR-1992 (Rel. 21, Created)
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Negative regulator of mitosis.
GN Name=BIME;
OS Emericella nidulans (Aspergillus nidulans).
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;
OC Eurotiales; Trichocomaceae; Emericella.
OX NCBI_TaxID=162425;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90375468; PubMed=1697851;
RA Engle D.B., Osmani S.A., Osmani A.H., Rosborough S., Xiang X.,
RA Morris N.R.;
RT "A negative regulator of mitosis in Aspergillus is a putative
membrane-spanning protein."
RL J. Biol. Chem. 265:16132-16137(1990).
CC -!- FUNCTION: Negative regulator of mitosis in E.nidulans. This
protein is part of a regulatory pathway that includes the nima
protein kinase. It is required to prevent premature entry into
mitosis. Mutations to this protein both cause cells to enter
mitosis and prevent them from leaving mitosis.
CC -!- SIMILARITY: Contains 4 PC repeats.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
between the Swiss Institute of Bioinformatics and the EMBL outstation -
the European Bioinformatics Institute. There are no restrictions on its
use by non-profit institutions as long as its content is in no way
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entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
or send an email to license@isb-sib.ch).
CC -----
CC EMBL; M59705; AAA51478.1; -.
DR PIR; A37879; BWASBE.
DR InterPro; IPR002015; APC_proteasome.
DR Pfam; PF01851; PC_rep; 4.
DR Mitosis; Repeat; Transmembrane.
KW DOMAIN 342 353 Nuclear localization signal (Potential).
FT TRANSMEM 1623 1643 Potential.
FT TRANSMEM 1685 1703 Potential.
FT TRANSMEM 1746 1764 Potential.
SQ SEQUENCE 2073 AA; 229178 MW; 05E4E81EADDF51E4 CRC64;

Query Match 77.8%; Score 42; DB 1; Length 2073;
Best Local Similarity 80.0%; Pred. No. 56;
Matches 8; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
Qy 1 TRLTRKRGK 10
|||||:
Db 832 TRLTRKRGK 841

RN [3]
 RP SEQUENCE FROM N.A., AND VARIANTS ILE-98 AND GLU-4181.
 RX MEDLINE=87008488; PubMed=3759943;
 RA Chen S.-H., Yang C.-Y., Chen P.-F., Setzer D., Tanimura M., Li W.-H.,
 RA Gatto A.M. Jr., Chan L.;
 RT "The complete cDNA and amino acid sequence of human apolipoprotein B-
 RT 100.";
 RL J. Biol. Chem. 261:12918-12921(1986).
 RN [4]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87041416; PubMed=3464946;
 RA Law S.W., Grant S.M., Higuchi K., Hospattankar A.V., Lackner K.J.,
 RA Lee N., Brewer H.B. Jr.;
 RT "Human liver apolipoprotein B-100 cDNA: complete nucleic acid and
 RT derived amino acid sequence.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:8142-8146(1986).
 RN [5]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=87161758; PubMed=3030729;
 RA Cladaras C., Hadzopoulou-Cladaras M., Nolte R.T., Atkinson D.,
 RA Zannis V.I.;
 RT "The complete sequence and structural analysis of human apolipoprotein
 RT B-100: relationship between apoB-100 and apoB-48 forms.";
 RL EMO J. 5:3495-3507(1986).
 RN [6]
 RP SEQUENCE OF 709-906 FROM N.A.
 RX MEDLINE=85270450; PubMed=3860836;
 RA Deeb S.S., Motulsky A.G., Albers J.J.;
 RT "A partial cDNA clone for human apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 82:4983-4986(1985).
 RN [7]
 RP SEQUENCE OF 3056-3159 FROM N.A.
 RX MEDLINE=86041888; PubMed=3903660;
 RA Mehriban M., Schumaker V.N., Fareed G.C., West R., Johnson D.F.,
 RA Kirchgesner T.G., Lin H.-C., Wang X., Ma Y., Mendiaz E., Lusa A.J.;
 RT "Human apolipoprotein B: identification of cDNA clones and
 RT characterization of mRNA.";
 RL Nucleic Acids Res. 13:6937-6953(1985).
 RN [8]
 RP SEQUENCE OF 1937-2018 AND 3811-4334 FROM N.A., AND VARIANT GLU-4181.
 RX MEDLINE=86093680; PubMed=3841204;
 RA Carlsson P., Olofsson S.O., Bondjers G., Darnfors C., Wiklund O.,
 RA Bjursell G.;
 RT "Molecular cloning of human apolipoprotein B cDNA.";
 RL Nucleic Acids Res. 13:8813-8826(1985).
 RN [9]
 RP SEQUENCE OF 3109-4563 FROM N.A.
 RX MEDLINE=85300528; PubMed=2994225;
 RA Knott T.J., Rall S.C. Jr., Innerarity T.L., Jacobson S.F., Urdea M.S.,
 RA Levy-Wilson B., Powell L.M., Pease R.J., Eddy R., Nakai H., Byers M.,
 RA Priestley L.M., Robertson E., Rall L.B., Betsholtz C., Shows T.B.,
 RA Mahley R.W., Scott J.;
 RT "Human apolipoprotein B: structure of carboxyl-terminal domains, sites
 RT of gene expression, and chromosomal localization.";
 RL Science 230:37-43(1985).
 RN [10]
 RP SEQUENCE OF 1-291 FROM N.A.
 RX MEDLINE=86149325; PubMed=3513177;
 RA Protter A.A., Hardman D.A., Schilling J.W., Miller J., Appleby V.,
 RA Chen G.C., Kirsher S.W., McEnroe G., Kane J.P.;
 RT "Isolation of a cDNA clone encoding the amino-terminal region of human
 RT apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:1467-1471(1986).
 RN [11]
 RP SEQUENCE OF 1-1670 FROM N.A., AND VARIANT ILE-98.
 RX MEDLINE=86287319; PubMed=3461454;
 RA Protter A.A., Hardman D.A., Sato K.Y., Schilling J.W., Yamanaka M.,
 RA Hort V.J., Hjertild K.A., Chen G.C., Kane J.P.;
 RT "Analysis of cDNA clones encoding the entire B-26 region of human
 RT apolipoprotein B.";
 RL Proc. Natl. Acad. Sci. U.S.A. 83:5678-5682(1986).
 RN [12]
 RP PARTIAL SEQUENCE, AND IDENTIFICATION OF APO-B48.
 RX

RX MEDLINE=88018019; PubMed=3659919;
 RA Chen S.-H., Habib G., Yang C.-H., Gu Z.-W., Lee B.R., Weng S.-H.,
 RA Silberman S.R., Cai S.-J., Deslypere J.P., Rosseneu M.,
 RA Gatto A.M. Jr., Li W.-H., Chan L.;
 RT "Apolipoprotein B-48 is the product of a messenger RNA with an organ-
 RT specific in-frame stop codon.";
 RL Science 238:363-366(1987).
 RN [13]
 RP DOMAINS.
 RX MEDLINE=87039351; PubMed=3773997;
 RA Knott T.C., Pease R.J., Powell L.M., Wallis S.C., Rall S.C. Jr.,
 RA Innerarity T.L., Blackhart B., Taylor W.R., Marcel Y., Milne R.,
 RA Johnson D., Fuller M., Lusa A.J., McCarthy B.J., Mahley R.W.,
 RA Levy-Wilson B., Scott J.;
 RT "Complete protein sequence and identification of structural domains of
 RT human apolipoprotein B.";
 RL Nature 323:734-738(1986).
 RN [14]
 RP DOMAINS.
 RA Yang C.-Y., Chen S.-H., Gianturco S.H., Bradley W.A., Sparrow J.T.,
 RA Tanimura M., Li W.-H., Sparrow D.A., Deloof H., Rosseneu M.,
 RA Lee F.-S., Gu Z.-W., Gatto A.M. Jr., Chan L.;
 RT "Sequence, structure, receptor-binding domains and internal repeats of
 RT human apolipoprotein B-100.";
 RL Nature 323:738-742(1986).
 RN [15]
 RP CALCIUM-BINDING DATA.
 RX MEDLINE=86242245; PubMed=3087360;
 RA Dashti N., Lee D.M., Mok T.;
 RT "Apolipoprotein B is a calcium binding protein.";
 RL Biochem. Biophys. Res. Commun. 137:493-499(1986).
 RN [16]
 RP PALMITOYLATION OF CVS-1112.
 RX MEDLINE=20143590; PubMed=10679026;
 RA Zhao Y., McCabe J.B., Vance J., Berthiaume L.G.;
 RT "Palmitoylation of apolipoprotein B is required for proper
 RT intracellular sorting and transport of cholesterol esters and
 RT triglycerides.";
 RL Mol. Biol. Cell 11:721-734(2000).
 RN [17]
 RP VARIANT SER-4338.
 RX MEDLINE=91071750; PubMed=1979313;
 RA Navajas M., Laurent A.-M., Moreel J.-F., Ragab A., Cambou J.-P.,
 RA Cuny G., Cambien F., Roizes G.;
 RT "Detection by denaturing gradient gel electrophoresis of a new
 RT polymorphism in the apolipoprotein B gene.";
 RL Hum. Genet. 86:91-93(1990).
 RN [18]
 RP VARIANT FDB GLN-3527.
 RX MEDLINE=89098975; PubMed=2563166;
 RA Soria L.F., Ludwig E.H., Clarke H.R.G., Vega G.L., Grundy S.M.,
 RA McCarthy B.J.;
 RT "Association between a specific apolipoprotein B mutation and familial
 RT defective apolipoprotein B-100.";
 RL Proc. Natl. Acad. Sci. U.S.A. 86:587-591(1989).
 RN [19]
 RP VARIANT LEU-2739.
 RX MEDLINE=91016974; PubMed=2216805;
 RA Huang L.-S., Gavish D., Breslow J.L.;
 RT "Sequence polymorphism in the human apoB gene at position 8344.";
 RL Nucleic Acids Res. 18:5922-5922(1990).
 RN [20]
 RP VARIANT FDB CYS-3558.
 RX MEDLINE=95190020; PubMed=7883971;
 RA Pullinger C.R., Hennessey L.K., Chatterton J.E., Liu W., Love J.A.,
 RA Mendel C.M., Frost P.H., Malloy M.J., Schumaker V.N., Kane J.P.;
 RT "Familial ligand-defective apolipoprotein B. Identification of a new
 RT mutation that decreases LDL receptor binding affinity.";
 RL J. Clin. Invest. 95:1225-1234(1995).
 RN [21]
 RP VARIANTS LEU-1437; SER-1914; LYS-2566; THR-3121; ALA-3945; MET-4128
 RP AND THR-4481.
 RX MEDLINE=97044521; PubMed=8889592;
 RX

RA Poixier O., Ricard S., Behague I., Souriau C., Evans A.E.,
 RA Arveiler D., Marques-Vidal P., Luc G., Roizes G., Cambien F.,
 RT "Detection of new variants in the apolipoprotein B (Apo B) gene by
 RT PCR-SSCP.",
 RL Hum. Mutat. 8:282-285 (1996).
 RN [22]
 RP VARIANTS FDB GLN-3527 AND CYS-3558.
 RX MEDLINE=97403938; PubMed=9259199;
 RA Rabes J.P., Varret M., Saint-Jore B., Erlich D., Jondeau G.,
 RA Krempf M., Giraudet P., Junien C., Boileau C.,
 RT "Familial ligand-defective apolipoprotein B-100: simultaneous
 RT detection of the ARG3500-->GLN and ARG3531-->CYS mutations in a French
 RT population.",
 RL Hum. Mutat. 10:160-163 (1997).
 RN [23]
 RP VARIANTS SER-1914; ARG-1923; LEU-2739; ASP-3319; THR-3427; GLN-3432
 RP AND ILE-3921.
 RX MEDLINE=98141125; PubMed=9490296;
 RA Leren T.P., Bakken K.S., Hoel V., Hjermann I., Berg K.,
 RT "Screening for mutations of the apolipoprotein B gene causing
 RT hypocholesterolemia.",
 RL Hum. Genet. 102:44-49 (1998).
 CC -!- FUNCTION: Apolipoprotein B is a major protein constituent of
 CC chylomicrons (apo B-48), LDL (apo B-100) and VLDL (apo B-100). Apo
 CC B-100 functions as a recognition signal for the cellular binding
 CC and internalization of LDL particles by the apoB/E receptor.
 CC -!- SUBCELLULAR LOCATION: Secreted.

Query Match 71.3%; Score 38.5; DB 1; Length 4563;
 Best Local Similarity 90.9%; Pred. No. 6.9e+02;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKRGGLK 11
 ||||| |||||

Db 3385 TRLTRK-RGLK 3394

RESULT 6

Q7Z600
 ID Q7Z600 PRELIMINARY; PRT; 4563 AA.
 AC Q7Z600;
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Apolipoprotein B (Including Ag(X) antigen).
 GN Name=APOB;
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OC NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
 RA Ahearn M.O., Kuldane S.A., Rajkumar N., Toth E.J., Yi Q.,
 RA Nickerson D.A.,
 RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
 RX EMBL; AY324608; AAF72970.1; -.
 DR GO; GO:0005319; F:lipid transporter activity; IEA.
 DR GO; GO:0006869; P:lipid transport; IEA.
 DR InterPro; IPR009454; DUF1081.
 DR InterPro; IPR001747; Lipid_transprt_N.
 DR Pfam; PF06448; DUF1081; 1.
 DR Pfam; PF01347; Vitellogenin_N; 1.
 DR SMART; SMO0638; LPD_N; 1.
 KW Lipoprotein.
 SQ SEQUENCE 4563 AA; 515553 MW; 030B34167CEBC63C CRC64;

Query Match 71.3%; Score 38.5; DB 2; Length 4563;
 Best Local Similarity 90.9%; Pred. No. 6.9e+02;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKRGGLK 11
 ||||| |||||

Db 3385 TRLTRK-RGLK 3394

RESULT 7

Q7U8V5
 ID Q7U8V5 PRELIMINARY; PRT; 431 AA.
 AC Q7U8V5;
 DT 01-OCT-2003 (TrEMBLrel. 25, Created)
 DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Possible GTPase.
 GN OrderedLocusNames=SYNW0504;
 OS Synechococcus sp. (strain WH8102).
 OC Bacteria; Cyanobacteria; Chroococcales; Synechococcus.
 OC NCBI_TaxID=84588;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=22825697; PubMed=12917641; DOI=10.1038/nature01943;
 RA Palenik B., Brahamsha B., Larimer F.W., Land M.L., Hauser L.,
 RA Chain P., Lamerdin J.E., Regala W., Allen E.E., McCarren J.,
 RA Paulsen I.T., Dufresne A., Partensky F., Webb E.A., Waterbury J.,
 RT "The genome of a motile marine Synechococcus.",
 RL Nature 424:1037-1042 (2003).
 RX EMBL; BX569690; CAE07019.1; -.
 DR GO; GO:0005525; F:GTP binding; IEA.
 DR InterPro; IPR005225; Small_GTP.
 DR TIGRFAMs; TIGR00231; small_GTP; 1.
 KW Complete proteome.
 SQ SEQUENCE 431 AA; 47647 MW; 666ECD647A83217D CRC64;

Query Match 68.5%; Score 37; DB 2; Length 431;
 Best Local Similarity 77.8%; Pred. No. 1e+02;
 Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

QY 2 RLTRKRGGL 10
 ||||| |||||

Db 27 RLTRREGGL 35

RESULT 8

Q9LXV0
 ID Q9LXV0 PRELIMINARY; PRT; 488 AA.
 AC Q9LXV0;
 DT 01-OCT-2000 (TrEMBLrel. 15, Created)
 DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
 DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
 DE Glucosyltransferase-like protein.
 GN Name=T24H18_60;
 OS Arabidopsis thaliana (Mouse-ear cress).
 OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
 OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
 OC eurosida II; Brassicales; Brassicaceae; Arabidopsia.
 OC NCBI_TaxID=3702;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Bevan M., Robben J., Grymonprez B., Volckaert G., Bancroft I.,
 RA Mewes H.W., Rudd S., Lemcke K., Mayer K.F.X.,
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE FROM N.A.
 RA EU Arabidopsis sequencing project;
 RL Submitted (APR-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- SIMILARITY: Belongs to the UDP-glucosyltransferase family.
 DR EMBL; AL353013; CAB88253.1; -.
 DR FIR; T49903; T49903.
 DR GO; GO:0016758; F:transferase activity, transferring hexosyl . . .; IEA.
 DR GO; GO:0008152; P:metabolism; IEA.
 DR InterPro; IPR002213; UDP_gluco_trans.
 DR Pfam; PF00201; UDPGT; 1.
 DR PROSITE; PS00375; UDPGT; 1.
 KW Glucosyltransferase; Transferase.
 SQ SEQUENCE 488 AA; 54867 MW; B6CC2E0A55452647 CRC64;


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Query Match      68.5%; Score 37; DB 2; Length 488;
Best Local Similarity 77.8%; Pred. No. 1.2e+02;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY      2 RLTRKERGL 10
DB      344 RLTRSERGL 352

RESULT 9
Q8VZF9 PRELIMINARY; PRT; 488 AA.
AC Q8VZF9;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE AT5g12890/T24H18.60.
OS Arabidopsis thaliana (Mouse-ear cress).
OC Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
OC Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids;
OC eurosids II; Brassicales; Brassicaceae; Arabidopsis.
OX NCBI_TaxID=3702;
RN [1]
RP SEQUENCE FROM N.A.
RA Cheuk R., Chen H., Kim C.J., Meyers M.C., Banh J., Bowser L.,
RA Carninci P., Chang E., Dale J.M., Goldsmith A.D., Hayashizaki Y.,
RA Ishida J., Jones T., Kamiya A., Karlin-Neumann G., Kawai J., Lam B.,
RA Lee J.M., Lin J., Miranda M., Narusaka M., Nguyen M., Onodera C.S.,
RA Palm C.J., Quach H.L., Sakurai T., Satou M., Seki M., Southwick A.,
RA Tang C.C., Toriumi M., Wu H.C., Yamada K., Yamamura Y., Yu G., Yu S.,
RA Shinozaki K., Davis R.W., Theologis A., Ecker J.R.;
RL Submitted (NOV-2001) to the EMBL/GenBank/DBJ databases.
CC -!- SIMILARITY: Belongs to the UDP-glycosyltransferase family.
DR GO; GO:0016758; F:transferase activity, transferring hexosyl . . .; IEA.
DR GO; GO:0008152; P:metabolism; IEA.
DR InterPro; IPR002213; UDP_glycos_trans.
DR Pfam; PF00201; UDPGT; 1.
DR PROSITE; PS00375; UDPGT; 1.
DR GlycoStltransferase; Transferase.
KW SEQUENCE 488 AA; 54793 MW; B6CC3E07FE389D37 CRC64;

Query Match      68.5%; Score 37; DB 2; Length 488;
Best Local Similarity 77.8%; Pred. No. 1.2e+02;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY      2 RLTRKERGL 10
DB      344 RLTRSERGL 352

RESULT 10
Q89I78 PRELIMINARY; PRT; 94 AA.
AC Q89I78;
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-JUN-2003 (TrEMBLrel. 24, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Bcl5761 protein.
GN OrderedLocNames=bcl5761;
OS Bradyrhizobium japonicum.
OC Bacteria; Proteobacteria; Alphaproteobacteria; Rhizobiales;
OC Bradyrhizobiaceae; Bradyrhizobium.
OX NCBI_TaxID=375;
RN [1]
RP SEQUENCE FROM N.A.
RA STRAIN=USDA110;
RX MEDLINE=22484998; PubMed=12597275;
RA Kaneko T., Nakamura Y., Sato S., Minamisawa K., Uchiumi T.,
RA Sasamoto S., Watanabe A., Ideawa K., Iriguchi M., Kawashima K.,
RA Kohara M., Matsumoto M., Shimpō S., Tsuruoka H., Wada T., Yamada M.,
RA Tabata S.;
RP "Complete genomic sequence of nitrogen-fixing symbiotic bacterium

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RT Bradyrhizobium japonicum USDA110.";
RL DNA Res. 9:189-197(2002).
DR EMBL; AP005956; BACS1026.1; -.
KW Complete proteome.
SQ SEQUENCE 94 AA; 10344 MW; 4925ED8E76A04BB8 CRC64;

Query Match      66.7%; Score 36; DB 2; Length 94;
Best Local Similarity 87.5%; Pred. No. 32;
Matches 7; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      2 RLTRKERG 9
DB      86 RLTRREERG 93

RESULT 11
Q6HM04 PRELIMINARY; PRT; 248 AA.
AC Q6HM04;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DE Hypothetical protein.
GN ORFNames=Bf9727_1080;
OS Bacillus thuringiensis serovar konkukian str. 97-27.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus;
OC Bacillus thuringiensis serovar konkukian.
OX NCBI_TaxID=281309;
RN [1]
RP SEQUENCE FROM N.A.
RA Brettn T.S., Bruce D., Challacombe J.F., Gilna P., Han C., Hill K.,
RA Hitchcock P., Jackson P., Keim P., Longmire J., Lucas S., Okinaka R.,
RA Richardson P., Rubin E., Tice H.,
RA Submitted (JUN-2004) to the EMBL/GenBank/DBJ databases.
RL EMBL; AE017355; AA763074.1; -.
DR InterPro; IPR010983; EF_Hand_like.
KW Hypothetical protein.
SQ SEQUENCE 248 AA; 30052 MW; 4C44DCDD6B421736 CRC64;

Query Match      66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 91;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY      3 LTRKERGLK 11
DB      155 LTRKERQLK 163

RESULT 12
Q73BX2 PRELIMINARY; PRT; 248 AA.
AC Q73BX2;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DE Hypothetical protein.
GN OrderedLocNames=BCE1296;
OS Bacillus cereus (Strain ATCC 10987).
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=222523;
RN [1]
RP SEQUENCE FROM N.A.
RX PubMed=14960714;
RA Raško D.A., Ravel J., Oekstad O.A., Helgason E., Cer R.Z., Jiang L.,
RA Shores K.A., Fouts D.E., Tourasse N.J., Angioli S.V., Kolonay J.F.,
RA Nelson W.C., Kolstoe A.-B., Fraser C.M., Read T.D.;
RT "The genome sequence of Bacillus cereus ATCC 10987 reveals metabolic
RT adaptations and a large plasmid related to Bacillus anthracis pXO1.";
RL Nucleic Acids Res. 32:977-988(2004).
DR EMBL; AE017268; AAS40225.1; -.
DR TIGR; BCE1296; -.
DR InterPro; IPR010983; EF_Hand_like.

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KW Complete proteome; Hypothetical protein.
SQ SEQUENCE 248 AA; 30064 MW; 2EC2A2C58C7CA2EA CRC64;

Query Match 66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 91;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 LTRKERGLK 11
Db 155 LTRKERQLK 163

RESULT 13
ID Q81GL8 PRELIMINARY; PRT; 248 AA.
AC Q81GL8;
DT 01-JUN-2003 (TRENBLrel. 24, Created)
DT 01-JUN-2003 (TRENBLrel. 24, Last sequence update)
DE Hypothetical protein.
GN ORFNames=BC1176;
OS Bacillus cereus (strain ATCC 14579 / DSM 31).
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=226900;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22608415; PubMed=12721630; DOI=10.1038/nature01582;
RA Ivanova N., Sorokin A., Anderson I., Galleron N., Candelon B.,
RA Kapatal V., Bhattacharya A., Reznik G., Mikhailova N., Lapidus A.,
RA Chu L., Mazur M., Goltsman E., Larsen N., D'Souza M., Walunas T.,
RA Grechkin Y., Pusch G., Haseikorn R., Fongstein M., Ehrlich S.D.,
RA Overbeek R., Kyrpides N.C.;
RT "Genome sequence of Bacillus cereus and comparative analysis with
RT Bacillus anthracis."
RL Nature 423:87-91(2003).
DR EMBL; AE017001; AAP08163.1; -
DR InterPro; IPR010983; EF_Hand_like.
KW Hypothetical protein.
SQ SEQUENCE 248 AA; 29788 MW; 79EBCC5746C2BDA3 CRC64;

Query Match 66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 91;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 LTRKERGLK 11
Db 155 LTRKERQLK 163

RESULT 14
ID Q81TS7 PRELIMINARY; PRT; 248 AA.
AC Q81TS7; Q61210; Q6KVU7;
DT 01-JUN-2003 (TRENBLrel. 24, Created)
DT 01-JUN-2003 (TRENBLrel. 24, Last sequence update)
DE Hypothetical protein.
GN OrderedLocusNames=BAL187, BAS1098; ORFNames=GBAA1187;
OS Bacillus anthracis.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=1392;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=Ames / Isolate Porton;
RA Read T.D., Peterson S.N., Tourasse N.J., Baillie L.W., Paulsen I.T.,
RA Nelson K.E., Tettelin H., Fouts D.E., Eisen J.A., Gill S.R.,
RA Holtzapple E.K., Okstad O.A., Helgason E., Ristone J., Wu M.,
RA Kolonay J.F., Beanan M.J., Dodson R.J., Brinkac L.M., Gwinn M.L.,
RA DeBoy R.T., Madupu R., Dougherty S.C., Durkin A.S., Haft D.H.,
RA Nelson W.C., Peterson J.D., Pop M., Khouri H.M., Radune D.,
RA Benton J.L., Mahmoud Y., Jiang L., Hance I.R., Weidman J.F.,
RA Berry K.J., Plaut R.D., Wolf A.M., Watkins K.L., Nierman W.C.,

HAZEN A., CLINE R.T., REDMOND C., THWAITE J.E., WHITE O.,
SAZBERG S.L., THOMASON B., FRIEDLANDER A.M., KOEHLER T.M.,
HANNA P.C., KOLSTOE A.-B., FRASER C.M.;
"THE GENOME SEQUENCE OF BACILLUS ANTHRACIS AMES AND COMPARISON TO
CLOSELY RELATED BACTERIA.";
RL Nature 423:81-86(2003).
RN [2]
RP SEQUENCE FROM N.A.
RX STRAIN=Ames / isolate 0581;
RA Ravel J., Rasko D.A., Shumway M.F., Jiang L., Cer R.Z., Federova N.B.,
RA Wilson M., Stanley S., Decker S., Read T.D., Salzberg S.L.,
RA Fraser C.M.;
RT "Bacillus anthracis comparative genomics.";
RL Submitted (MAY-2004) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE FROM N.A.
RX STRAIN=Sterne;
RA Brettin T.S., Bruce D., Challacombe J.F., Gilna P., Han C., Hill K.,
RA Hitchcock P., Jackson P., Keim P., Longmire J., Lucas S., Okinaka R.,
RA Richardson P., Rubin E., Tice H.;
RL Submitted (JAN-2004) to the EMBL/GenBank/DBJ databases.
DR EMBL; AE017027; AAP25150.1; -
DR EMBL; AE017334; AAT30275.1; -
DR EMBL; AE017225; AAT53421.1; -
DR TIGR; BA1187; -
DR InterPro; IPR010983; EF_Hand_like.
KW Hypothetical protein.
SQ SEQUENCE 248 AA; 30052 MW; 4C44DCDD6B421736 CRC64;

Query Match 66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 91;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 LTRKERGLK 11
Db 155 LTRKERQLK 163

RESULT 15
ID Q9K8Y0 PRELIMINARY; PRT; 248 AA.
AC Q9K8Y0;
DT 01-OCT-2000 (TRENBLrel. 15, Created)
DT 01-OCT-2000 (TRENBLrel. 15, Last sequence update)
DE BH2872 protein.
GN Name=BH2872;
OS Bacillus halodurans.
OC Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.
OX NCBI_TaxID=86665;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=C-125;
RX MEDLINE=20512582; PubMed=11058132;
RA Takami H., Nakasone K., Takaki Y., Maeno G., Sasaki R., Masui N.,
RA Fuji F., Hirama C., Nakamura Y., Ogasawara N., Kuhara S.,
RA Horikoshi K.;
RT "Complete genome sequence of the alkaliphilic bacterium Bacillus
RT halodurans and genomic sequence comparison with Bacillus subtilis.";
RL Nucleic Acids Res. 28:4317-4331(2000).
DR EMBL; AP001516; BAB06591.1; -
DR PIR; H84008; H84008.
SQ SEQUENCE 248 AA; 29874 MW; C75106082942A7DE CRC64;

Query Match 66.7%; Score 36; DB 2; Length 248;
Best Local Similarity 88.9%; Pred. No. 91;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 LTRKERGLK 11
Db 155 LTRKERQLK 163

```

Search completed: January 13, 2005, 01:51:08
Job time : 88.8361 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:14:40 ; Search time 84.9344 Seconds
(without alignments)
46.460 Million cell updates/sec

Title: US-09-823-418-19

Perfect score: 55

Sequence: 1 TRLTRKDRGLK 11

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 2002273 seqs, 358729299 residues

Total number of hits satisfying chosen parameters: 2002273

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : A_Geneseq_23Sep04:*

1: Geneseqp1980s:*

2: Geneseqp1990s:*

3: Geneseqp2000s:*

4: Geneseqp2001s:*

5: Geneseqp2002s:*

6: Geneseqp2003as:*

7: Geneseqp2003bs:*

8: Geneseqp2004s:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB | ID | Description |
|------------|-------|-------------|--------|----|----------|--------------------|
| 1 | 55 | 100.0 | 11 | 2 | AAY30700 | Aay30700 Apo-B100 |
| 2 | 51 | 92.7 | 11 | 2 | AAY30698 | Aay30698 Apo-B100 |
| 3 | 44 | 80.0 | 11 | 2 | AAY30697 | Aay30697 Apo-B100 |
| 4 | 42 | 76.4 | 11 | 2 | AAY30699 | Aay30699 Apo-B100 |
| 5 | 39.5 | 71.8 | 10 | 2 | AAY30683 | Aay30683 Apo-B100 |
| 6 | 39.5 | 71.8 | 10 | 2 | AAY30691 | Aay30691 Apo-B100 |
| 7 | 39 | 70.9 | 621 | 6 | ABU24550 | Protein e |
| 8 | 39 | 70.9 | 1756 | 8 | ADN47108 | Adn47108 Thermococ |
| 9 | 38.5 | 70.0 | 11 | 2 | AAW57205 | Aaw57205 Apo B bin |
| 10 | 38.5 | 70.0 | 13 | 2 | AAW57207 | Aaw57207 Apo B 100 |
| 11 | 38.5 | 70.0 | 15 | 2 | AAW41261 | Aaw41261 Apolipop |
| 12 | 38.5 | 70.0 | 15 | 2 | AAW96892 | Aaw96892 ApoB-100 |
| 13 | 38.5 | 70.0 | 20 | 6 | ABJ37575 | Abj37575 Heparin b |
| 14 | 38.5 | 70.0 | 22 | 2 | AAW57208 | Aaw57208 Apo B 100 |
| 15 | 38.5 | 70.0 | 22 | 2 | AAW57209 | Aaw57209 Apo B 100 |
| 16 | 38.5 | 70.0 | 34 | 5 | AAE14541 | Aae14541 Human apo |
| 17 | 38.5 | 70.0 | 36 | 2 | AAW96876 | Aaw96876 Nucleic a |
| 18 | 38.5 | 70.0 | 37 | 2 | AAW64587 | Aaw64587 Human apo |
| 19 | 38.5 | 70.0 | 51 | 2 | AAW96845 | Aaw96845 Nucleic a |
| 20 | 38.5 | 70.0 | 343 | 4 | ABB37687 | Abb37687 Peptide # |
| 21 | 38.5 | 70.0 | 343 | 4 | ABG52504 | Abg52504 Human liv |
| 22 | 38.5 | 70.0 | 377 | 2 | AAW72704 | Aaw72704 Human apo |
| 23 | 38.5 | 70.0 | 377 | 2 | AAW34031 | Aaw34031 Sequence |
| 24 | 38.5 | 70.0 | 2463 | 8 | ADJ57400 | Adj57400 Human apo |
| 25 | 38.5 | 70.0 | 3923 | 2 | AAY31237 | Aay31237 Human Apo |

ALIGNMENTS

RESULT 1

AAV30700

ID AAY30700 standard; peptide; 11 AA.

XX AC AAY30700;

XX XX 17-NOV-1999 (first entry)

XX DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.

XX KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;

XX KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.

XX OS Synthetic.

XX OS Homo sapiens.

XX PN W09946598-A1.

XX PD 16-SEP-1999.

XX PF 05-MAR-1999; 99WO-US004805.

XX PR 10-MAR-1998; 98US-0077618P.

XX PA (REGC) UNIV CALIFORNIA.

XX PI Innerarity TL, Boren JOS;

XX DR WPI; 1999-551509/46.

XX PT Identifying compounds which affect binding of low density lipoprotein with proteoglycan, used for, e.g. obtaining compounds for reducing atherosclerosis.

XX PS Claim 17; Page 58; 70pp; English.

XX CC AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan receptor mutations. They were created to identify compounds which modulate atherosclerosis. The peptides are derived from amino acids 3358 to 3367 of apoB100. The method comprises detecting compounds which affect low density lipoprotein (LDL) binding with proteoglycan (PG). The method can be used for identifying compounds which disrupt LDL-PG binding without inhibiting LDL receptor binding. Such compounds can be used to reduce or prevent the formation of atherosclerotic lesions and prevent atherosclerosis. The transgenic non-human animals and mammals which express human apo-B100 can be used as an in vivo model system for the study of atherosclerosis, and in vivo assay methods for identifying compounds which modulate atherosclerosis and/or LDL-PG binding. They can

CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX

SQ Sequence 11 AA;
 Query Match 100.0%; Score 55; DB 2; Length 11;
 Best Local Similarity 100.0%; Pred. No. 0.0007;
 Matches 11; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKDRGLK 11
 |||||:||||
 Db 1 TRLTRKDRGLK 11

RESULT 2
 AAY30698
 ID AAY30698 standard; peptide; 11 AA.
 AC AAY30698;
 XX
 DT 17-NOV-1999 (first entry)
 XX
 DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
 XX

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN WO9946598-A1.
 XX
 PD 16-SEP-1999.

XX 05-MAR-1999; 99WO-US004805.
 XX 10-MAR-1998; 98US-0077618P.
 XX (REGC) UNIV CALIFORNIA.
 XX Innerarity TL, Boren JOS;
 XX WPI; 1999-551509/46.
 XX

Identifying compounds which affect binding of low density lipoprotein
 with proteoglycan, used for, e.g. obtaining compounds for reducing
 atherosclerosis.

Claim 17; Page 57; 70pp; English.

AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 receptor mutations. They were created to identify compounds which
 modulate atherosclerosis. The peptides are derived from amino acids 3358
 to 3367 of apoB100. The method comprises detecting compounds which affect
 low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 can be used for identifying compounds which disrupt LDL-PG binding
 without inhibiting LDL receptor binding. Such compounds can be used to
 reduce or prevent the formation of atherosclerotic lesions and prevent
 atherosclerosis. The transgenic non-human animals and mammals which
 express human apo-B100 can be used as an in vivo model system for the
 study of atherosclerosis, and in vivo assay methods for identifying
 compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 also be used to identify compounds which result in an increase in
 atherosclerotic regions. Thus the assays may be used to determine whether
 a particular food or drug composition tends to stimulate or inhibit the
 formation of atherosclerotic lesions. The polynucleotides can also be
 used in gene therapy for preventing or reducing the severity of
 atherosclerosis in an animal or mammal

SQ Sequence 11 AA;

Query Match 92.7%; Score 51; DB 2; Length 11;
 Best Local Similarity 90.9%; Pred. No. 0.0043;
 Matches 10; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKDRGLK 11
 |||||:||||
 Db 1 TRLTRKDRGLK 11

RESULT 3
 AAY30697
 ID AAY30697 standard; peptide; 11 AA.
 AC AAY30697;
 XX

XX 17-NOV-1999 (first entry)
 XX
 DE Apo-B100 derived peptide showing a proteoglycan receptor mutation.
 XX

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN WO9946598-A1.
 XX
 PD 16-SEP-1999.

XX 05-MAR-1999; 99WO-US004805.
 XX 10-MAR-1998; 98US-0077618P.
 XX (REGC) UNIV CALIFORNIA.
 XX Innerarity TL, Boren JOS;
 XX WPI; 1999-551509/46.

Identifying compounds which affect binding of low density lipoprotein
 with proteoglycan, used for, e.g. obtaining compounds for reducing
 atherosclerosis.
 Claim 17; Page 57; 70pp; English.
 AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 receptor mutations. They were created to identify compounds which
 modulate atherosclerosis. The peptides are derived from amino acids 3358
 to 3367 of apoB100. The method comprises detecting compounds which affect
 low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 can be used for identifying compounds which disrupt LDL-PG binding
 without inhibiting LDL receptor binding. Such compounds can be used to
 reduce or prevent the formation of atherosclerotic lesions and prevent
 atherosclerosis. The transgenic non-human animals and mammals which
 express human apo-B100 can be used as an in vivo model system for the
 study of atherosclerosis, and in vivo assay methods for identifying
 compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 also be used to identify compounds which result in an increase in
 atherosclerotic regions. Thus the assays may be used to determine whether
 a particular food or drug composition tends to stimulate or inhibit the
 formation of atherosclerotic lesions. The polynucleotides can also be
 used in gene therapy for preventing or reducing the severity of
 atherosclerosis in an animal or mammal

Sequence 11 AA;

Query Match 80.0%; Score 44; DB 2; Length 11;
 Best Local Similarity 81.8%; Pred. No. 0.1;
 Matches 9; Conservative 1; Mismatches 1; Indels 1; Gaps 0;

QY 1 TRLTRKDRGLK 11

```
Db          1 TRLTRKRGK 11
|||||: ||||
RESULT 4
AAAY30699
ID  AAAY30699 standard; peptide; 11 AA.
XX
AC  AAAY30699;
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW  Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW  low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS  Synthetic.
OS  Homo sapiens.
XX
PN  WO9946598-A1.
XX
PD  16-SEP-1999.
XX
PF  05-MAR-1999; 99WO-US004805.
XX
PR  10-MAR-1998; 98US-0077618P.
XX
PA  (REGC ) UNIV CALIFORNIA.
XX
PI  Innerarity TL, Boren JOS;
XX
WPI; 1999-551509/46.
XX
PT  Identifying compounds which affect binding of low density lipoprotein
PT  with proteoglycan, used for, e.g. obtaining compounds for reducing
PT  atherosclerosis.
XX
PS  Claim 17; Page 58; 70pp; English.
XX
CC  AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC  receptor mutations. They were created to identify compounds which
CC  modulate atherosclerosis. The peptides are derived from amino acids 3358
CC  to 3367 of apoB100. The method comprises detecting compounds which affect
CC  low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC  can be used for identifying compounds which disrupt LDL-PG binding
CC  without inhibiting LDL receptor binding. Such compounds can be used to
CC  reduce or prevent the formation of atherosclerotic lesions and prevent
CC  atherosclerosis. The transgenic non-human animals and mammals which
CC  express human apo-B100 can be used as an in vivo model system for the
CC  study of atherosclerosis, and in vivo assay methods for identifying
CC  compounds which modulate atherosclerosis and/or LDL-PG binding. They can
CC  also be used to identify compounds which result in an increase in
CC  atherosclerotic regions. Thus the assays may be used to determine whether
CC  a particular food or drug composition tends to stimulate or inhibit the
CC  formation of atherosclerotic lesions. The polynucleotides can also be
CC  used in gene therapy for preventing or reducing the severity of
CC  atherosclerosis in an animal or mammal
XX
SQ  Sequence 11 AA;
Query Match 76.4%; Score 42; DB 2; Length 11;
Best Local Similarity 81.8%; Pred. No. 0.26;
Matches 9; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy          1 TRLTRKRGK 11
|||||: ||||
Db          1 TRLTRKRGK 11
|||||: ||||
RESULT 5
AAAY30683
ID  AAAY30683 standard; peptide; 10 AA.
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
KW  Apo-B100; proteoglycan receptor mutation; atherosclerosis;
KW  low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
XX
OS  Synthetic.
OS  Homo sapiens.
XX
PN  WO9946598-A1.
XX
PD  16-SEP-1999.
XX
PF  05-MAR-1999; 99WO-US004805.
XX
PR  10-MAR-1998; 98US-0077618P.
XX
PA  (REGC ) UNIV CALIFORNIA.
XX
PI  Innerarity TL, Boren JOS;
XX
WPI; 1999-551509/46.
XX
PT  Identifying compounds which affect binding of low density lipoprotein
PT  with proteoglycan, used for, e.g. obtaining compounds for reducing
PT  atherosclerosis.
XX
PS  Claim 17; Page 57; 70pp; English.
XX
CC  AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
CC  receptor mutations. They were created to identify compounds which
CC  modulate atherosclerosis. The peptides are derived from amino acids 3358
CC  to 3367 of apoB100. The method comprises detecting compounds which affect
CC  low density lipoprotein (LDL) binding with proteoglycan (PG). The method
CC  can be used for identifying compounds which disrupt LDL-PG binding
CC  without inhibiting LDL receptor binding. Such compounds can be used to
CC  reduce or prevent the formation of atherosclerotic lesions and prevent
CC  atherosclerosis. The transgenic non-human animals and mammals which
CC  express human apo-B100 can be used as an in vivo model system for the
CC  study of atherosclerosis, and in vivo assay methods for identifying
CC  compounds which modulate atherosclerosis and/or LDL-PG binding. They can
CC  also be used to identify compounds which result in an increase in
CC  atherosclerotic regions. Thus the assays may be used to determine whether
CC  a particular food or drug composition tends to stimulate or inhibit the
CC  formation of atherosclerotic lesions. The polynucleotides can also be
CC  used in gene therapy for preventing or reducing the severity of
CC  atherosclerosis in an animal or mammal
XX
SQ  Sequence 10 AA;
Query Match 71.8%; Score 39.5; DB 2; Length 10;
Best Local Similarity 90.3%; Pred. No. 0.73;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
Qy          1 TRLTRKRGK 11
|||||: ||||
Db          1 TRLTR-DRGLK 10
|||||: ||||
RESULT 6
AAAY30691
ID  AAAY30691 standard; peptide; 10 AA.
XX
AC  AAAY30691;
XX
DT  17-NOV-1999 (first entry)
XX
DE  Apo-B100 derived peptide showing a proteoglycan receptor mutation.
XX
```

KW Apo-B100; proteoglycan receptor mutation; atherosclerosis;
 KW low density lipoprotein; proteoglycan; LDL; atherosclerotic lesion.
 XX Synthetic.
 OS Homo sapiens.
 XX W09946598-A1.
 PN W09946598-A1.
 XX 16-SEP-1999.
 PD 05-MAR-1999; 99WO-US004805.
 XX 10-MAR-1998; 98US-0077618P.
 PR (REGC) UNIV CALIFORNIA.
 PA Innerarity TL, Boren JOS;
 PI WPI; 1999-551509/46.
 XX Identifying compounds which affect binding of low density lipoprotein
 PT with proteoglycan, used for, e.g. obtaining compounds for reducing
 PT atherosclerosis.
 XX Claim 17; Page 57; 70pp; English.
 PS AAY30582-Y30700 represent apo-B100 derived peptides showing proteoglycan
 CC receptor mutations. They were created to identify compounds which
 CC modulate atherosclerosis. The peptides are derived from amino acids 3358
 CC to 3367 of apoB100. The method comprises detecting compounds which affect
 CC low density lipoprotein (LDL) binding with proteoglycan (PG). The method
 CC can be used for identifying compounds which disrupt LDL-PG binding
 CC without inhibiting LDL receptor binding. Such compounds can be used to
 CC reduce or prevent the formation of atherosclerotic lesions and prevent
 CC atherosclerosis. The transgenic non-human animals and mammals which
 CC express human apo-B100 can be used as an in vivo model system for the
 CC study of atherosclerosis, and in vivo assay methods for identifying
 CC compounds which modulate atherosclerosis and/or LDL-PG binding. They can
 CC also be used to identify compounds which result in an increase in
 CC atherosclerotic regions. Thus the assays may be used to determine whether
 CC a particular food or drug composition tends to stimulate or inhibit the
 CC formation of atherosclerotic lesions. The polynucleotides can also be
 CC used in gene therapy for preventing or reducing the severity of
 CC atherosclerosis in an animal or mammal
 XX Sequence 10 AA;
 SQ
 Query Match 71.8%; Score 39.5; DB 2; Length 10;
 Best Local Similarity 90.9%; Pred. No. 0.73;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRKDRGLK 11
 Db 1 TRLTRKDRGLK 11
 |||||
 1 TRLTRKDRGLK 11
 RESULT 7
 ABU24550
 ID ABU24550 standard; protein; 621 AA.
 XX
 AC ABU24550;
 XX
 XX 19-JUN-2003 (first entry)
 DT Protein encoded by Prokaryotic essential gene #10077.
 DE Antisense; prokaryotic essential gene; cell proliferation; drug design.
 XX Clostridium botulinum.
 OS W020027183-A2.
 PN W020027183-A2.
 XX 03-OCT-2002.
 PD

XX 21-MAR-2002; 2002WO-US009107.
 PF 21-MAR-2001; 2001US-00815242.
 PR 06-SEP-2001; 2001US-00948993.
 PR 25-OCT-2001; 2001US-0342923P.
 PR 08-FEB-2002; 2002US-00072851.
 PR 06-MAR-2002; 2002US-0362699P.
 XX (ELIT-) ELITRA PHARM INC.
 PA Wang L, Zamudio C, Malone C, Haselbeck R, Ohlsen KL, Zyskind JW;
 PI Wall D, Trawick JD, Carr GJ, Yamamoto R, Forsyth RA, Xu HH;
 XX WPI; 2003-029926/02.
 DR N-PSDB; ACA28420.
 DR New antisense nucleic acids, useful for identifying proteins or screening
 PT for homologous nucleic acids required for cellular proliferation to
 PT isolate candidate molecules for rational drug discovery programs.
 XX Claim 25; SEQ ID NO 52474; 1766pp; English.
 XX The invention relates to an isolated nucleic acid comprising any one of
 CC the 6213 antisense sequences given in the specification where expression
 CC of the nucleic acid inhibits proliferation of a cell. Also included are:
 CC (1) a vector comprising a promoter operably linked to the nucleic acid
 CC encoding a polypeptide whose expression is inhibited by the antisense
 CC nucleic acid; (2) a host cell containing the vector; (3) an isolated
 CC polypeptide or its fragment whose expression is inhibited by the
 CC antisense nucleic acid; (4) an antibody capable of specifically binding
 CC the polypeptide; (5) producing the polypeptide; (6) inhibiting cellular
 CC proliferation or the activity of a gene in an operon required for
 CC proliferation; (7) identifying a compound that influences the activity of
 CC the gene product or that has an activity against a biological pathway
 CC required for proliferation, or that inhibits cellular proliferation; (8)
 CC identifying a gene required for cellular proliferation or the biological
 CC pathway in which a proliferation-required gene or its gene product lies
 CC or a gene on which the test compound that inhibits proliferation of an
 CC organism acts; (9) manufacturing an antibiotic; (10) profiling a
 CC compound's activity; (11) a culture comprising strains in which the gene
 CC product is overexpressed or underexpressed; (12) determining the extent
 CC to which each of the strains is present in a culture or collection of
 CC strains; or (13) identifying the target of a compound that inhibits the
 CC proliferation of an organism. The antisense nucleic acids are useful for
 CC identifying proteins or screening for homologous nucleic acids required
 CC for cellular proliferation to isolate candidate molecules for rational
 CC drug discovery programs, or for screening homologous nucleic acids
 CC required for proliferation in cells other than *S. aureus*, *S. typhimurium*,
 CC *K. pneumoniae* or *P. aeruginosa*. The present sequence is encoded by one of
 CC the target prokaryotic essential genes. Note: the sequence data for this
 CC patent did not form part of the printed specification, but was obtained
 CC in electronic format directly from WIPO at
 CC ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 621 AA;
 Query Match 70.9%; Score 39; DB 6; Length 621;
 Best Local Similarity 77.8%; Pred. No. 68;
 Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
 QY 2 RLTRKDRGL 10
 Db 228 RLTRKDRGV 236
 |||||
 RESULT 8
 ADN47108
 ID ADN47108 standard; protein; 1756 AA.
 XX
 AC ADN47108;
 XX 01-JUL-2004 (first entry)
 DT

XX Thermococcus kodakaraensis KOD1 protein sequence SeqID986.
 DE
 XX
 KW gene disruption; gene targeting; marker gene; transformation;
 KW homologous recombination; hyperthermostable archaeobacterium; KOD1;
 KW gene structure; gene function; enzyme activity; medicine;
 KW forensic science; food; drug inspection; molecular biology; immunology.
 XX
 OS Thermococcus kodakaraensis.
 XX WO2004022736-A1.
 XX 18-MAR-2004.
 XX
 XX 29-AUG-2003; 2003WO-IB003597.
 XX
 XX 30-AUG-2002; 2002JP-00319011.
 XX
 XX (NISC-) JAPAN SCI & TECHNOLOGY CORP.
 XX
 XX Imanaka T, Atomi H;
 XX WPI; 2004-257583/24.
 XX
 XX Method for disrupting targeted gene in genome of organism particularly
 PT thermostable bacterium and with genome chips for analysis, applicable in
 PT studying gene structure and functions.
 XX
 XX Claim 9; SEQ ID NO 986; 598pp; Japanese.
 XX
 XX This invention relates to a novel method for targeting disruption of an
 CC arbitrary gene in a genome of an organism which comprises providing the
 CC whole sequential data of the genome of such organism, selecting at least
 CC 1 arbitrary region in the sequence, providing a vector that contains a
 CC sequence homologous with the selected region and a marker gene.
 CC transformation, and homologous recombination. The genome is preferably
 CC the genome of a hyperthermostable archaeobacterium, particularly
 CC Thermococcus kodakaraensis KOD1. The method is for targeting the
 CC disruption of a gene in the genome of an organism, which is applicable in
 CC studying gene structure and functions as well as enzyme activities of
 CC encoded proteins and useful in medicine, forensic science, food or drug
 CC inspection, molecular biology and immunology. With this method, the
 CC disruption of a gene at an arbitrary position in a genome can be achieved
 CC efficiently and reliably. The present sequence is that of a protein
 CC encoded by the genome of Thermococcus kodakaraensis which was derived
 CC using the method of the invention. Note: The sequence data for this
 CC patent did not form part of the printed specification, but was obtained
 CC in electronic format directly from WIPO at
 XX ftp.wipo.int/pub/published_pct_sequences
 XX
 SQ Sequence 1756 AA;
 Query Match 70.9%; Score 39; DB 8; Length 1756;
 Best Local Similarity 88.9%; Pred. No. 2e+02;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 2 RLTRKDRGL 10
 Db 1183 RLTSKDRGL 1191
 RESULT 9
 AAW57205
 ID AAW57205 standard; peptide; 11 AA.
 XX
 AC AAW57205;
 XX
 XX 03-AUG-1998 (first entry)
 DT
 XX Apo B binding site peptide 2.
 DE
 XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW

KW LDL; receptor component; apo B100 receptor site.
 XX Synthetic.
 OS
 XX WO9813385-A2.
 PN
 XX 02-APR-1998.
 PD
 XX 25-SEP-1997; 97WO-GB002610.
 XX
 XX 27-SEP-1996; 96GB-00020153.
 PR
 XX (UYST) UNIV STRATHCLYDE.
 FA
 XX Halbert GW, Owens MD, Baillie G;
 PI WPI; 1998-230637/20.
 XX
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 XX Claim 12; Page 52; 73pp; English.
 PS
 XX The present sequence represents a specifically claimed Apo B binding site
 CC peptide which can be used as a component of a non-naturally occurring,
 CC receptor-competent low density lipoprotein (LDL) particle of the present
 CC invention. The LDL particle comprises at least 1 peptide component that
 CC has at least 1 binding site for an apo B protein receptor and at least 1
 CC lipophilic substituent. Also described in the invention are peptides
 CC containing an apo B binding sequence with at least 70% identity with
 CC sequences: KAEYKKNRHH (1) or TRLTRKRGK (2), or their dimers. Non-
 CC naturally occurring, receptor-competent LDL particles are useful as: (i)
 CC drug-targeting vectors for delivering anticancer drugs to cancer cells
 CC that express an apo B protein receptor, and (ii) additives for cell
 CC culture media especially as growth supplements. Non-naturally occurring,
 CC receptor-competent LDL particles do not require the complete apo B
 CC sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 11 AA;
 Query Match 70.0%; Score 38.5; DB 2; Length 11;
 Best Local Similarity 90.9%; Pred. No. 1.3;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 Qy 1 TRLTRKDRGLK 11
 Db 2 TRLTRK-RGLK 11
 RESULT 10
 AAW57207
 ID AAW57207 standard; peptide; 13 AA.
 XX
 AC AAW57207;
 XX
 XX 03-AUG-1998 (first entry)
 DT
 XX Apo B 100 binding site peptide analogue peptide B.
 DE
 XX Apo B; binding site; receptor; cancer; drug delivery; anticancer;
 KW growth supplement; non-natural lipid particle; low density lipoprotein;
 KW LDL; receptor component; apo B100 receptor site.
 XX
 XX Synthetic.
 OS
 XX Key Location/Qualifiers
 FH Modified-site 1
 FT /note= "attached to retinoic acid"
 XX
 XX WO9813385-A2.
 PN
 XX

PD 02-APR-1998.
 XX
 PF 25-SEP-1997; 97WO-GB002610.
 XX
 PR 27-SEP-1996; 96GB-00020153.
 XX
 PA (UYST) UNIV STRATHCLYDE.
 XX
 PI Halbert GW, Owens MD, Baillie G;
 XX WPI; 1998-230637/20.
 DR
 XX Non-natural lipid particle comprising peptide binding to apo B protein
 PT receptor - useful as, e.g. vector for delivering drugs to cancer cells
 PT that express this receptor.
 XX
 XX Claim 13; Fig 7; 73pp; English.
 XX
 CC The present sequence represents a specifically claimed Apo B 100 binding
 CC site peptide analogue which can be used as a component of a non-
 CC naturally occurring, receptor-competent low density lipoprotein (LDL)
 CC particle of the present invention. The LDL particle comprises at least 1
 CC peptide component that has at least 1 binding site for an apo B protein
 CC receptor and at least 1 lipophilic substituent. Also described in the
 CC invention are peptides containing an apo B binding sequence with at least
 CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRGK (2), or their
 CC dimers. Non-naturally occurring, receptor-competent LDL particles are
 CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to
 CC cancer cells that express an apo B protein receptor, and (ii) additives
 CC for cell culture media especially as growth supplements. Non-naturally
 CC occurring, receptor-competent LDL particles do not require the complete
 CC apo B sequence, which is large and tends to aggregate, to provide binding
 CC affinity to an apo B protein receptor
 XX
 SQ Sequence 13 AA;
 Query Match 70.0%; Score 38.5; DB 2; Length 13;
 Best Local Similarity 90.9%; Pred. No. 1.5;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRKRGK 11
 Db 3 TRLTRK-RGLK 12
 RESULT 11
 AAW41261
 ID AAW41261 standard; peptide; 15 AA.
 XX
 AC AAW41261;
 XX
 DT 19-MAY-1998 (first entry)
 XX
 DE Apolipoprotein B-100 fragment.
 XX
 KW Anti-coagulant; apolipoprotein B-100; apoB-100; metastatic spread;
 KW thromboplastin-mediated process; cancer; inhibitor; blood coagulation;
 KW angiogenesis; cellular differentiation; apoptosis; KRAD-14;
 KW prothrombinase complex.
 XX
 OS Synthetic.
 OS Homo sapiens.
 XX
 PN WO9743311-A1.
 XX
 PD 20-NOV-1997.
 XX
 PF 09-MAY-1997; 97WO-GB001255.
 XX
 PR 09-MAY-1996; 96GB-00009702.
 XX
 XX (UNLO) ROYAL FREE HOSPITAL SCHOOL MED.
 PA
 XX
 PI Bruckdorfer KR, Ettelaie C;
 XX WPI; 1998-008798/01.
 XX
 PT Peptide fragments of apo:apo:protein B-100 with anticoagulant activity -
 PT used for treating or preventing coagulation, inhibiting angiogenesis,
 PT cell differentiation and apoptosis.
 XX
 XX Disclosure; Page 22; 60pp; English.
 PS
 XX This sequence is an example of the peptide of the invention. It has the
 CC formula (I), or their variants with one or more internal deletions,
 CC insertions or substitutions, while retaining anti-coagulant properties of
 CC apolipoprotein B-100 (apoB-100). Z1-KAQ-X1-KKNKRRHS-X2-T-22 (I) X1 = S or
 CC Y; X2 = T or I; Z1 = the N terminus of the peptide, or 1-47 amino acids
 CC (aa); Z2 = the C terminus of the peptide, a terminal amide group or 1-77
 CC aa. Compositions containing the peptide are used for simultaneous,
 CC separate or sequential treatment of cancer, particularly to prevent
 CC metastatic spread. They are also used to inhibit thromboplastin-mediated
 CC processes, specifically to prevent or reduce blood coagulation (e.g.
 CC during or after surgery or in cases of heart attack, stroke etc.) and to
 CC inhibit angiogenesis, cellular differentiation or apoptosis. KRAD-14,
 CC which is active as such or as part of a 98-aa peptide, inhibits
 CC activation of the prothrombinase complex; and prevents activation of
 CC factor VII on the surface of thromboplastin and of platelets by thrombin.
 CC It binds to the residues 58-66 of thromboplastin. Since (I) are much
 CC smaller than apoB-100, they act more quickly
 XX
 SQ Sequence 15 AA;
 Query Match 70.0%; Score 38.5; DB 2; Length 15;
 Best Local Similarity 90.9%; Pred. No. 1.7;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 QY 1 TRLTRKRGK 11
 Db 1 TRLTRK-RGLK 10
 RESULT 12
 AAW96892
 ID AAW96892 standard; peptide; 15 AA.
 XX
 AC AAW96892;
 XX
 DT 22-APR-1999 (first entry)
 XX
 DE ApoB-100 nuclear localisation signal sequence, residues 3353-3367.
 XX
 KW Human apolipoprotein B-100; apoB-100; very-low density lipoprotein; VLDL;
 KW apolipoprotein; binding; in vivo transport; nucleic acid; binding domain;
 KW nuclear localisation sequence; gene therapy; cancer; cystic fibrosis;
 KW non-small cell lung carcinoma; diabetes; arteriosclerosis.
 XX
 OS Homo sapiens.
 OS
 PN WO9856938-A1.
 XX
 PD 17-DEC-1998.
 XX
 PF 10-JUN-1998; 98WO-US011927.
 XX
 PR 13-JUN-1997; 97US-00874807.
 PR 14-MAY-1998; 98US-00079030.
 XX
 XX (BAYU) BAYLOR COLLEGE MEDICINE.
 XX
 PI Guevara JG, Hoogveen RC, Moore JP;
 XX WPI; 1999-070331/06.
 DR
 XX Composition comprising nucleic acid bound to LDL or VLDL lipoprotein -
 PT used for delivering nucleic acid to cells for gene therapy and antisense

PT treatment.

XX Claim 19; Fig 13D; 293pp; English.

PS AAW96878-97 represent nuclear localisation signal sequence derived from

CC human apolipoprotein B-100 (apoB-100). ApoB-100 is a major apoprotein

CC component of very-low density lipoproteins (VLDL), intermediate density

CC lipoprotein (IDL), low density lipoproteins (LDL) and lipoprotein a. The

CC present sequence can be used in the composition of the invention. The

CC specification describes a composition that comprises LDL and

CC apolipoproteins for the binding and in vivo transport of nucleic acids.

CC The composition is used to deliver nucleic acids to eukaryotic cells, in

CC vivo or in vitro, for expressing a therapeutic polypeptide or antisense

CC molecule (or ribozyme). Specifically they are used for gene therapy of

CC cancers (particularly non-small cell lung carcinoma), diabetes, cystic

CC fibrosis and arteriosclerosis

XX

SQ Sequence 15 AA;

Query Match 70.0%; Score 38.5; DB 2; Length 15;

Best Local Similarity 90.9%; Pred. No. 1.7;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKDRGLK 11

Db ||||| |||||

6 TRLTRK-RGLK 15

RESULT 13

ABU37575

ID ABU37575 standard; peptide; 20 AA.

XX

AC ABU37575;

XX

DT 10-MAY-2003 (first entry)

XX

DE Heparin binding peptide sequence #28.

XX

KW Cytostatic; antirheumatic; antiarthritic; antidiabetic; ophthalmological;

KW cardiovascular; circulatory; ligand; sulphated; sulphonated; tumour;

KW rheumatoid arthritis; hypoxia; diabetic retinopathy; heparin.

XX

OS Unidentified.

XX

PN WO2003007689-A2.

XX

PD 30-JAN-2003.

XX

PF 22-JUL-2002; 2002WO-US023419.

XX

PR 20-JUL-2001; 2001US-0306726P.

XX

PA (ETHZ-) ETH ZUERICH.

PA (UYZU-) UNIV ZURICH.

XX

PI Hubbell JA, Schoenmakers R, Maynard HD;

XX

DR WPI; 2003-300420/29.

XX

PT Use of a ligand comprising of at least one sulfated or sulfonated amino

PT acid for the treatment of e.g. tumors, rheumatoid arthritis, diabetic

PT retinopathy and hypoxia.

XX

PS Disclosure; Fig 2; 79pp; English.

XX

CC The invention relates to a novel ligand for binding a target biomolecule,

CC which comprises a peptide having at least one sulphated or sulphonated

CC amino acid and at least one amino acid chosen from neutral and positively

CC charged amino acids. The novel ligands can be used for the treatment of

CC e.g. tumours, rheumatoid arthritis, diabetic retinopathy and hypoxia.

CC This sequence represents a heparin binding peptide relating to the

CC invention

SQ Sequence 20 AA;

Query Match 70.0%; Score 38.5; DB 6; Length 20;

Best Local Similarity 90.9%; Pred. No. 2.4;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKDRGLK 11

Db ||||| |||||

7 TRLTRK-RGLK 16

RESULT 14

AAW57208

ID AAW57208 standard; peptide; 22 AA.

XX

AC AAW57208;

XX

DT 03-AUG-1998 (first entry)

XX

DE Apo B 100 binding site peptide analogue peptide C.

XX

KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;

KW growth supplement; non-natural lipid particle; low density lipoprotein;

KW LDL; receptor component; apo B100 receptor site.

XX

OS Synthetic.

XX

FH Key Location/Qualifiers

FT Modified-site 1 /note= "attached to retinoic acid"

FT Modified-site 22 /note= "attached to cholesterol"

XX

PN WO9813385-A2.

XX

PD 02-APR-1998.

XX

PF 25-SEP-1997; 97WO-GB002610.

XX

PR 27-SEP-1996; 96GB-00020153.

XX

PA (UYST) UNIV STRATHCLYDE.

XX

PI Halbert GW, Owens MD, Baillie G;

XX

DR WPI; 1998-230637/20.

XX

PT Non-natural lipid particle comprising peptide binding to apo B protein

PT receptor - useful as, e.g. vector for delivering drugs to cancer cells

PT that express this receptor.

XX

PS Claim 13; Fig 7; 73pp; English.

XX

CC The present sequence represents a specifically claimed Apo B 100 binding

CC site peptide analogue which can be used as a component of a non-

CC naturally occurring, receptor-competent low density lipoprotein (LDL)

CC particle of the present invention. The LDL particle comprises at least 1

CC peptide component that has at least 1 binding site for an apo B protein

CC receptor and at least 1 lipophilic substituent. Also described in the

CC invention are peptides containing an apo B binding sequence with at least

CC 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKRGGLK (2), or their

CC dimers. Non-naturally occurring, receptor-competent LDL particles are

CC useful as: (i) drug-targeting vectors for delivering anticancer drugs to

CC cancer cells that express an apo B protein receptor, and (ii) additives

CC for cell culture media especially as growth supplements. Non-naturally

CC occurring, receptor-competent LDL particles do not require the complete

CC apo B sequence, which is large and tends to aggregate, to provide binding

CC affinity to an apo B protein receptor

XX

SQ Sequence 22 AA;

Query Match 70.0%; Score 38.5; DB 2; Length 22;

Best Local Similarity 90.9%; Pred. No. 2.6;

Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
QY 1 TRLTRKDRGLK 11
 |||||
Db 7 TRLTRK-RGLK 16

Search completed: January 13, 2005, 01:43:05
Job time : 86.1011 secs

RESULT 15
AAW57209
ID AAW57209 standard; peptide; 22 AA.
XX
AC AAW57209;
XX
DT 03-AUG-1998 (first entry)
XX
DE Apo B 100 binding site peptide analogue peptide D.
XX
KW Apo B; binding site; receptor; cancer; drug delivery; anticancer;
KW growth supplement; non-natural lipid particle; low density lipoprotein;
KW LDL; receptor component; apo B100 receptor site.
XX
OS Synthetic.

FH Key Location/Qualifiers
FT Modified-site 1
FT /note= "attached to retinoic acid"

PN W09813385-A2.
XX
PD 02-APR-1998.
XX
XX 25-SEP-1997; 97WO-GB002610.
XX 27-SEP-1996; 96GB-00020153.
XX (UYST) UNIV STRATHCLYDE.
XX
XX Halbert GW, Owens MD, Baillie G;
XX WPI; 1998-230637/20.
XX
XX Non-natural lipid particle comprising peptide binding to apo B protein
XX receptor - useful as, e.g. vector for delivering drugs to cancer cells
XX that express this receptor.

PS Claim 13; Fig 7; 73pp; English.
XX
XX The present sequence represents a specifically claimed Apo B 100 binding
XX site peptide analogue which can be used as a component of a non-
XX naturally occurring, receptor-competent low density lipoprotein (LDL)
XX particle of the present invention. The LDL particle comprises at least 1
XX peptide component that has at least 1 binding site for an apo B protein
XX receptor and at least 1 lipophilic substituent. Also described in the
XX invention are peptides containing an apo B binding sequence with at least
XX 70% identity with sequences: KAEYKKNKRRH (1) or TRLTRKGLK (2), or their
XX dimers. Non-naturally occurring, receptor-competent LDL particles are
XX useful as: (i) drug-targeting vectors for delivering anticancer drugs to
XX cancer cells that express an apo B protein receptor, and (ii) additives
XX for cell culture media especially as growth supplements. Non-naturally
XX occurring, receptor-competent LDL particles do not require the complete
XX apo B sequence, which is large and tends to aggregate, to provide binding
XX affinity to an apo B protein receptor

SQ Sequence 22 AA;
Query Match 70.0%; Score 38.5; DB 2; Length 22;
Best Local Similarity 90.9%; Pred. No. 2.6;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

QY 1 TRLTRKDRGLK 11
 |||||
Db 7 TRLTRK-RGLK 16

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OM protein - protein search, using sw model

Run On: January 13, 2005, 01:30:05 ; Search time 15.8689 Seconds
(without alignments)
66.696 Million cell updates/sec

Title: US-09-823-418-19
Perfect score: 55
Sequence: 1 TRLTRKDRGLK 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283416 seqs, 96216763 residues

Total number of hits satisfying chosen parameters: 283416

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 79:.*
1: pir1:.*
2: pir2:.*
3: pir3:.*
4: pir4:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query % | | Length | DB | ID | Description |
|------------|-------|---------|--------|--------|--------|----|--------------------|
| | | Match | Length | | | | |
| 1 | 40 | 72.7 | 250 | 2 | D98082 | | hypothetical prote |
| 2 | 40 | 72.7 | 2073 | 1 | BWASBE | | bimE protein - Eme |
| 3 | 39 | 70.9 | 807 | 2 | AC1031 | | hypothetical prote |
| 4 | 38.5 | 70.0 | 596 | 2 | S32802 | | apolipoprotein B - |
| 5 | 38.5 | 70.0 | 4563 | 1 | LPHUB | | apolipoprotein B-1 |
| 6 | 36 | 65.5 | 250 | 2 | F95218 | | hypothetical prote |
| 7 | 35 | 63.6 | 78 | 2 | B81098 | | hypothetical prote |
| 8 | 35 | 63.6 | 236 | 2 | AI2432 | | hypothetical prote |
| 9 | 35 | 63.6 | 321 | 2 | A37842 | | hypothetical prote |
| 10 | 35 | 63.6 | 384 | 2 | AS1987 | | hypothetical prote |
| 11 | 34.5 | 62.7 | 269 | 2 | C60950 | | apolipoprotein B-1 |
| 12 | 34.5 | 62.7 | 779 | 2 | JH0102 | | apolipoprotein B - |
| 13 | 34 | 61.8 | 195 | 2 | F71164 | | hypothetical prote |
| 14 | 34 | 61.8 | 331 | 2 | S73019 | | daunorubicin resis |
| 15 | 34 | 61.8 | 389 | 2 | D64337 | | 16S rRNA 5'-region |
| 16 | 34 | 61.8 | 405 | 2 | C72305 | | transposase, IS605 |
| 17 | 34 | 61.8 | 419 | 2 | A81651 | | conserved hypothet |
| 18 | 34 | 61.8 | 458 | 2 | E91092 | | probable invasion |
| 19 | 34 | 61.8 | 458 | 2 | A85938 | | probable invasion |
| 20 | 34 | 61.8 | 488 | 2 | T49903 | | Glucosyltransferas |
| 21 | 34 | 61.8 | 543 | 2 | F96624 | | hypothetical prote |
| 22 | 33 | 60.0 | 246 | 2 | C97177 | | C-terminal domain |
| 23 | 33 | 60.0 | 248 | 2 | H84008 | | hypothetical prote |
| 24 | 33 | 60.0 | 264 | 2 | A12112 | | hypothetical prote |
| 25 | 33 | 60.0 | 289 | 2 | T01531 | | hypothetical prote |
| 26 | 33 | 60.0 | 305 | 1 | NKVLHR | | core antigen - her |
| 27 | 33 | 60.0 | 341 | 2 | AC1148 | | hypothetical prote |
| 28 | 33 | 60.0 | 389 | 2 | AD0722 | | glutamate dehydrog |
| 29 | 33 | 60.0 | 419 | 2 | G84799 | | hypothetical prote |

| | | | | | | |
|----|------|------|------|---|--------|--------------------|
| 30 | 33 | 60.0 | 429 | 2 | H71491 | hypothetical prote |
| 31 | 33 | 60.0 | 503 | 2 | H86146 | hypothetical prote |
| 32 | 33 | 60.0 | 826 | 2 | E96720 | probable alpha-amy |
| 33 | 33 | 60.0 | 1451 | 2 | I40325 | dermonecrotic toxi |
| 34 | 33 | 60.0 | 3187 | 2 | JC5837 | 364K Golgi complex |
| 35 | 32.5 | 59.1 | 275 | 2 | E60950 | apolipoprotein B-1 |
| 36 | 32 | 58.2 | 100 | 2 | A36950 | urease (EC 3.5.1.5 |
| 37 | 32 | 58.2 | 107 | 2 | B85356 | glycine-rich prote |
| 38 | 32 | 58.2 | 196 | 2 | S69732 | hypothetical prote |
| 39 | 32 | 58.2 | 290 | 2 | H72071 | ct386 hypothetical |
| 40 | 32 | 58.2 | 290 | 2 | E86551 | CT386 hypothetical |
| 41 | 32 | 58.2 | 299 | 2 | D83010 | probable binding p |
| 42 | 32 | 58.2 | 300 | 2 | A81597 | conserved hypothet |
| 43 | 32 | 58.2 | 326 | 2 | JQ0855 | hypothetical 36.8K |
| 44 | 32 | 58.2 | 346 | 2 | S49963 | hypothetical prote |
| 45 | 32 | 58.2 | 398 | 2 | S40752 | hypothetical prote |

ALIGNMENTS

RESULT 1

D98082
hypothetical protein fecB [imported] - Streptococcus pneumoniae (strain R6)
C:Species: Streptococcus pneumoniae
C>Date: 22-Oct-2001 #sequence_revision 22-Oct-2001 #text_change 09-Jul-2004
C:Accession: D98082
R:Hoekings, J.A.; Alborn Jr., W.; Arnold, J.; Blaszczyk, L.; Burgett, S.; DeHoff, B.S.;
e, R.; LeBlanc, D.J.; Lee, L.N.; Lefkowitz, E.J.; Lu, J.; Matsushima, P.; McAhren, S.;
Y, P.; Sun, P.M.; Winkler, M.E.
J. Bacteriol. 183, 5709-5717, 2001
A:Authors: Yang, Y.; Young-Bellido, M.; Zhao, G.; Zook, C.; Baltz, R.H.; Jaskunas, S.R
A:Title: Genome of the Bacterium Streptococcus pneumoniae Strain R6.
A:Reference number: A97872; MUID:21429245; PMID:11544234
A:Accession: D98082
A>Status: preliminary
A:Molecule type: DNA
A:Residues: 1-250 <KUR>
A:Cross-references: UNIPROT:Q8DNJ3; GB:A8007317; PIDN:AAL00489.1; PID:gl5459361; GSPDB
C:Genetics:
A:Gene: fecB
C:Superfamily: inner membrane protein malK; ATP-binding cassette homology

Query Match 72.7% Score 40; DB 2; Length 250;
Best Local Similarity 70.0%; Pred. No. 2.8;
Matches 7; Conservative 3; Mismatches 0; Indels 0; Gaps 0;

Qy 1 TRLTRKDRGL 10
:|||||:
Db 47 SRLTKKDRGV 56

RESULT 2

BWASBE
bimE protein - Emericella nidulans
C:Species: Emericella nidulans, Aspergillus nidulans
C>Date: 31-Dec-1991 #sequence_revision 31-Dec-1991 #text_change 09-Jul-2004
C:Accession: A37879
R:Engle, D.B.; Omani, S.A.; Omani, A.H.; Rosborough, S.; Xiang, X.; Morris, N.R.
J. Biol. Chem. 265, 16132-16137, 1990
A:Title: A negative regulator of mitosis in Aspergillus is a putative membrane-spanning
A:Reference number: A37879; MUID:90375468; PMID:1697851
A:Accession: A37879
A:Molecule type: mRNA
A:Residues: 1-2073 <ENG>
A:Cross-references: UNIPROT:P24686; GB:M59705; GB:J05607; NID:gi68026; PIDN:AAA51478.1;
A>Note: In addition to three predicted transmembrane domains, there are several potent
asein kinase, and one sequence that resembles a nuclear localization signal
C:Comment: This protein is part of a regulatory pathway that includes the nimA protein
ter mitosis and prevent them from leaving mitosis.
C:Genetics:
A:Gene: bimE
C:Superfamily: bimE protein

C;Keywords: cell cycle control; mitosis; transmembrane protein
 F;1623-1643/Domain: transmembrane #status predicted <TM1>
 F;1685-1703/Domain: transmembrane #status predicted <TM2>
 F;1746-1764/Domain: transmembrane #status predicted <TM3>

Query Match 72.7%; Score 40; DB 1; Length 2073;
 Best Local Similarity 80.0%; Pred. No. 23;
 Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDRGL 10
 |||||: |||
 Db 832 TRLTRKDRGL 841

RESULT 3
 AC1031
 hypotheical protein STY4573 [imported] - Salmonella enterica subsp. enterica serovar Typhimurium
 C;Species: Salmonella enterica subsp. enterica serovar Typhi
 F;1685-1703/Domain: transmembrane #status predicted <TM2>
 C;Note: this species has also been called Salmonella typhi
 C;Date: 09-Nov-2001 #sequence_revision 09-Nov-2001 #text_change 18-Nov-2002
 C;Accession: AC1031
 R;Parkhill, J.; Dougan, G.; James, K.D.; Thomson, N.R.; Pickard, D.; Wain, J.; Churcher, T.; Connor, P.; Cronin, A.; Davis, P.; Davies, R.M.; Dowd, L.; White, N.; Farrar, S.; Moule, S.; O'Garra, P.
 Nature 413, 848-852, 2001
 A;Authors: Parry, C.; Quail, M.; Rutherford, K.; Simmonds, M.; Skelton, J.; Stevens, K.; A;Title: Complete genome sequence of a multiple drug resistant Salmonella enterica serovar Typhimurium
 A;Reference number: AB0502; MUID:21534947; PMID:11677608
 A;Accession: AC1031
 A;Status: preliminary
 A;Molecule type: DNA
 A;Residues: 1-807 <PAR>
 A;Title: Complete genome sequence of a multiple drug resistant Salmonella enterica serovar Typhimurium
 A;Cross-references: GB:AL513382; PIDN:CAD09348.1; PID:g16505348; GSPDB:GN00176
 C;Genetics:
 A;Gene: STY4573

Query Match 70.9%; Score 39; DB 2; Length 807;
 Best Local Similarity 88.9%; Pred. No. 14;
 Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
 Qy 2 RLTRKDRGL 10
 |||||: |||
 Db 508 RLTRADRG 516

RESULT 4
 S32802
 apolipoprotein B - crab-eating macaque (fragment)
 C;Species: Macaca fascicularis (crab-eating macaque)
 C;Date: 06-Jan-1995 #sequence_revision 06-Jan-1995 #text_change 09-Jul-2004
 C;Accession: S32802
 R;Pape, M.E.; Castle, C.K.; Murray, R.W.; Funk, G.M.; Hunt, C.E.; Marotti, K.R.; Melchior, B.; et al.
 Biochim. Biophys. Acta 1086, 326-334, 1991
 A;Title: Apo B metabolism in the cynomolgus monkey: evidence for post-transcriptional regulation of apo B mRNA
 A;Reference number: S32802; MUID:92075708; PMID:1742325
 A;Accession: S32802
 A;Status: preliminary
 A;Molecule type: mRNA
 A;Residues: 1-596 <PAR>
 A;Cross-references: UNIPROT:Q28473; EMBL:X15737; PIDN:CAA33755.1; PID:g93012
 C;Superfamily: apolipoprotein B

Query Match 70.0%; Score 38.5; DB 2; Length 596;
 Best Local Similarity 90.9%; Pred. No. 13;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 Qy 1 TRLTRKDRGLK 11
 |||||: |||
 Db 226 TRLTRK-RGLK 235

RESULT 5
 LPHUB

apolipoprotein B-100 precursor - human
 N;Contains: apolipoprotein B-26; apolipoprotein B-48; apolipoprotein B-74
 C;Species: Homo sapiens (man)
 C;Date: 28-Dec-1987 #sequence_revision 28-Dec-1987 #text_change 09-Jul-2004
 C;Accession: A27850; A25263; A25266; A25267; A25268; A25269; A25270; A25271; A25272; A25273; A25274; A25275; A25276; A25277; A25278; A25279; A25280; A25281; A25282; A25283; A25284; A25285; A25286; A25287; A25288; A25289; A25290; A25291; A25292; A25293; A25294; A25295; A25296; A25297; A25298; A25299; A25300; A25301; A25302; A25303; A25304; A25305; A25306; A25307; A25308; A25309; A25310; A25311; A25312; A25313; A25314; A25315; A25316; A25317; A25318; A25319; A25320; A25321; A25322; A25323; A25324; A25325; A25326; A25327; A25328; A25329; A25330; A25331; A25332; A25333; A25334; A25335; A25336; A25337; A25338; A25339; A25340; A25341; A25342; A25343; A25344; A25345; A25346; A25347; A25348; A25349; A25350; A25351; A25352; A25353; A25354; A25355; A25356; A25357; A25358; A25359; A25360; A25361; A25362; A25363; A25364; A25365; A25366; A25367; A25368; A25369; A25370; A25371; A25372; A25373; A25374; A25375; A25376; A25377; A25378; A25379; A25380; A25381; A25382; A25383; A25384; A25385; A25386; A25387; 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 A>Title: A partial cDNA clone for human apolipoprotein B.
 A;Reference number: A25774; MUID:85270450; PMID:3860836
 A;Accession: A25774
 A:Molecule type: mRNA
 A;Residues: 703-791, 'SSSWKAASHGCPHAGD', 810-906 <DE>
 A;Cross-references: GB:K03175; NID:g178821; PIDN:AAA51759.1; PID:g178822
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 A>Title: Analysis of the human apolipoprotein B gene: complete structure of the B-74 region
 A;Reference number: A31565; MUID:87191999; PMID:2883086
 A;Accession: A26533
 A:Molecule type: mRNA
 A;Residues: 1282-2721, 2742-3290, 'L', 3292-3336, 'N', 3338-3948, 'F', 3950-3963, 'Y', 3965-4180,
 A;Cross-references: GB:M15421; NID:g178817; PIDN:AAA51758.1; PID:g178818
 R;Hardman, D.A.; Protter, A.A.; Chen, G.C.; Schilling, J.W.; Sato, K.Y.; Lau, K.; Yamana
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 A>Title: Structural comparison of human apolipoproteins B-48 and B-100.
 A;Reference number: A29671; MUID:88050832; PMID:3676265
 A;Accession: A29671
 A:Molecule type: mRNA
 A;Residues: 1671-2323, 'PYW', 2327-2352, 'H', 2354-2398 <HAR>
 A;Cross-references: GB:M17367; NID:g178731; PIDN:AAA51741.1; PID:g178732
 R;Shoulders, C.C.; Myant, N.B.; Sidoli, A.; Rodriguez, J.C.; Cortese, C.; Baralle, F.E.;
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 A>Title: Molecular cloning of human LDL apolipoprotein B cDNA. Evidence for more than one
 A;Reference number: A90084; MUID:86130855; PMID:3841481
 A;Accession: A29287
 A:Molecule type: mRNA
 A;Residues: 3846-4298 <SHO>
 R;Pfitzner, R.; Wagener, R.; Stoffel, W.
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 A>Title: Isolation, expression and characterization of a human apolipoprotein B 100-spec
 A;Reference number: A25572; MUID:87076044; PMID:3024665
 A;Accession: A25572
 A:Molecule type: mRNA
 A;Residues: 4219-4337, 'S', 4339-4563 <PFI>
 A;Cross-references: GB:M36676
 R;Wei, C.F.; Chen, S.H.; Yang, C.Y.; Yang, C.Y.; Marcel, Y.L.; Milne, R.W.; Li, W.H.; Sparrow, J.T.;
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 A;Reference number: A24738; MUID:86042646; PMID:2933736
 A;Accession: A24738
 A:Molecule type: mRNA
 A;Residues: 'N', 3729-3731, 'I', 3733-3875, 'A', 3877-3948, 'F', 3950-3963, 'Y', 3965-3982, 'S', 39
 A;Cross-references: GB:M12413; NID:g178735; PIDN:AAA51742.1; PID:g178736
 R;Chen, S.H.; Habbib, G.; Yang, C.Y.; Gu, Z.W.; Lee, B.R.; Weng, S.; Silberman, S.R.; Cai
 Science 238, 363-366, 1987
 A>Title: Apolipoprotein B-48 is the product of a messenger RNA with an organ-specific in
 A;Reference number: A40133; MUID:88018019; PMID:3659919
 A;Accession: B40133
 A:Molecule type: mRNA
 A;Residues: 2155-2179 <CHI>
 A;Cross-references: GB:M18036; NID:g178799; PIDN:AAA51754.1; PID:g178800
 A;Note: this mRNA includes the stop codon of the organ-specific mRNA for apo48
 A;Accession: A40133
 A:Molecule type: protein
 A;Residues: 51-75, 101-110, 129-139, 158-174, 197-207, 276-287, 298-304, 306-314, 526-532, 538-55
 36; 1486-1498; 1537-1556; 1563-1572; 1601-1610; 1647-1661; 1697-1724; 1770-1781; 1859-1897; 1968-
 A;Note: these fragments were derived from apo48
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 A>Title: Carboxyl terminal analysis of human B-48 protein confirms the novel mechanism p
 A;Reference number: A28002; MUID:88106542; PMID:3426612
 A;Accession: A28002
 A:Molecule type: mRNA
 A;Residues: 2129-2179, 2181-2235 <HA2>
 A;Cross-references: GB:M18471
 A;Experimental source: intestine
 A;Note: this mRNA from intestine includes a stop codon created by RNA editing in place o
 R;Menraban, M.; Schumaker, V.N.; Fareed, G.C.; West, R.; Johnson, D.F.; Kirchgessner, T
 Nucleic Acids Res. 13, 6937-6953, 1985
 A>Title: Human apolipoprotein B: identification of cDNA clones and characterization of m

A;Reference number: A24269; MUID:86041888; PMID:3903660
 A;Accession: A24269
 A:Molecule type: mRNA
 A;Residues: 3056-3159 <MEH>
 A;Cross-references: GB:X03045; NID:g28783; PIDN:CAA26850.1; PID:g929609
 R;Hospatankar, A.V.; Higuchi, K.; Law, S.W.; Meglin, N.; Brewer Jr., H.B.
 Biochem. Biophys. Res. Commun. 148, 279-285, 1987
 A>Title: Identification of a novel in-frame translational stop codon in human intestine
 A;Reference number: A29659; MUID:88049670; PMID:2445342
 A;Accession: A29659
 A:Molecule type: mRNA
 A;Residues: 2169-2179 <HOS>
 A;Note: the sequence shown represents the carboxyl end of apolipoprotein B-48
 A;Note: two RNA species, 14.1kb in length, were isolated from the human intest
 ch encodes the 250K apoB-48, CAA encoding 2180-gln is substituted by the stop codon TAA
 R;Yang, C.; Kim, T.W.; Weng, S.; Lee, B.; Yang, M.; Gotto Jr., A.M.
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 A>Title: Isolation and characterization of sulfhydryl and disulfide peptides of human a
 A;Reference number: A35783; MUID:90319144; PMID:2115173
 A;Contents: disulfide bonds
 A;Accession: A35783
 A:Molecule type: protein
 A;Residues: 28-41, 76-97, 'I', 99-100; 175-193; 206-215; 239-249; 259-266; 357-399; 455-490; 512-
 A;Note: cysteines at positions 1112, 1422, 1505, 1662, 3761, 3917, and 4217 have free s
 R;LeBoeuf, R.C.; Miller, C.; Shively, J.E.; Schumaker, V.N.; Balla, M.A.; Lusis, A.J.
 FEBS Lett. 170, 105-108, 1984
 A>Title: Human apolipoprotein B: partial amino acid sequence.
 A;Reference number: A22006; MUID:84208786; PMID:6373369
 A;Accession: A22006
 A:Molecule type: protein
 A;Residues: 873-892, 'K', 894-896 <LE1>
 A;Accession: B22006
 A:Molecule type: protein
 A;Residues: 3113, 'L', 3115-3130, 'R', 3132-3133, 'P', 3135-3136, 'R' <LE2>
 R;Blackhart, B.D.; Ludwig, E.M.; Pierotti, V.R.; Caiati, L.; Onasch, M.A.; Wallis S.C
 J. Biol. Chem. 261, 15364-15367, 1986
 A>Title: Structure of the human apolipoprotein B gene.
 A;Reference number: A92564; MUID:87057153; PMID:2946672
 A;Contents: annotation; gene structure
 R;Wagener, R.; Pfitzner, R.; Stoffel, W.
 Biol. Chem. Hoppe-Seyler 368, 419-425, 1987
 A>Title: Studies on the organization of the human apolipoprotein B 100 gene.
 A;Reference number: A90715; MUID:87271140; PMID:2886136
 A;Contents: annotation; gene structure
 R;Weisgraber, K.H.; Rall Jr., S.C
 J. Biol. Chem. 262, 11097-11103, 1987
 A>Title: Human apolipoprotein B-100 heparin-binding sites.
 A;Reference number: A92605; MUID:87280197; PMID:3301850
 A;Contents: annotation; heparin binding and disulfide bond
 R;Dashti, N.; Lee, D.M.; Mok, T.
 Biochem. Biophys. Res. Commun. 137, 493-499, 1986
 A>Title: Apolipoprotein B is a calcium binding protein.
 A;Reference number: A90125; MUID:86242245; PMID:3087360
 A;Contents: annotation; calcium binding
 R;Carlsson, P.; Olofsson, S.O.; Bondjers, G.; Darnfors, C.; Wiklund, O.; Bjursell, G.
 Nucleic Acids Res. 13, 8813-8826, 1985
 A>Title: Molecular cloning of human apolipoprotein B cDNA.
 A;Reference number: 137178; MUID:86093680; PMID:3841204
 A;Accession: 137180

Query Match 70.0% Score 38.5; DB 1; Length 4563;
 Best Local Similarity 90.9%; Pred. No. 98;
 Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;
 Qy 1 TLTRKDRGLK 11
 ||||| |||||
 Db 3385 TLTRK-RGLK 3394

RESULT 6

F95218

hypothetical protein SPI871 [imported] - Streptococcus pneumoniae (strain TIGR4)

C;Species: Streptococcus pneumoniae
C;Date: 03-Aug-2001 #sequence_revision 03-Aug-2001 #text_change 09-Jul-2004
A;Accession: F95218
R;Tettelin, H.; Nelson, K.E.; Paulsen, I.T.; Eisen, J.A.; Read, T.D.; Peterson, S.; Heid
on, J.D.; Umayam, L.A.; White, O.; Salzberg, S.L.; Lewis, M.R.; Radune, D.; Holtzapple,
son, T.; Hickey, E.K.; Holt, I.E.
Science 293, 498-506, 2001
A;Authors: Loftus, B.J.; Yang, F.; Smith, H.O.; Venter, J.C.; Dougherty, B.A.; Morrison,
A;Title: Complete Genome Sequence of a virulent isolate of Streptococcus pneumoniae.
A;Reference number: A95000; MUID:21357209; PMID:11463916
A;Accession: F95218
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-250 <KUR>
A;Cross-references: UNIPROT:Q97NV0; GB:AE005672; PIDN:AAK75943.1; PID:g14973374; GSPDB:G
A;Experimental source: strain TIGR4
C;Genetics:
A;Gene: SPI871
C;Superfamily: inner membrane protein malk; ATP-binding cassette homology

Query Match 65.5%; Score 36; DB 2; Length 250;
Best Local Similarity 60.0%; Pred. No. 18;
Matches 6; Conservative 4; Mismatches 0; Indels 0; Gaps 0;

QY 1 TRLTRKDRGL 10
:|||||:
Db 47 SRLTKKQGV 56

RESULT 7
E81098
hypothetical protein NMB1311 [imported] - Neisseria meningitidis (strain MC58 serogroup
C;Species: Neisseria meningitidis
C;Date: 31-Mar-2000 #sequence_revision 31-Mar-2000 #text_change 09-Jul-2004
C;Accession: E81098
R;Tettelin, H.; Saunders, N.J.; Heidelberg, J.; Jeffries, A.C.; Nelson, K.E.; Eisen, J.A
Hickey, E.K.; Haft, D.H.; Salzberg, S.L.; White, O.; Fleischmann, R.D.; Dougherty, B.A.;
ri, H.; Qin, H.; Vamathevan, J.; Gill, J.; Scarlato, V.; Massignani, V.; Pizza, M.
Science 287, 1809-1815, 2000
A;Authors: Grandi, G.; Sun, L.; Smith, H.O.; Fraser, C.M.; Moxon, E.R.; Rappuoli, R.; Ve
A;Title: Complete genome sequence of Neisseria meningitidis serogroup B strain MC58.
A;Reference number: A81000; MUID:20175755; PMID:10710307
A;Accession: E81098
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-78 <TET>
A;Cross-references: UNIPROT:Q9JZ39; GB:AE002479; GB:AE002098; NID:g7226543; PIDN:AAF4168
A;Experimental source: serogroup B, strain MC58
C;Genetics:
A;Gene: NMB1311

Query Match 63.6%; Score 35; DB 2; Length 78;
Best Local Similarity 70.0%; Pred. No. 9;
Matches 7; Conservative 1; Mismatches 2; Indels 0; Gaps 0;

QY 2 RLTRKDRGLK 11
:|||||:
Db 26 RLTRKDRGRK 35

RESULT 8
AI2432
hypothetical protein all5017 [imported] - Nostoc sp. (strain PCC 7120)
C;Species: Nostoc sp. PCC 7120
A;Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C;Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Jul-2004
C;Accession: AI2432
R;Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iriguichi
Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata, S
DNA Res. 8, 205-213, 2001
A;Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium Ana
A;Reference number: AB1807; MUID:21595285; PMID:11759840
A;Accession: AI2432

A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-236 <KUR>
A;Cross-references: UNIPROT:Q8YMB7; GB:BA000019; PIDN:BAW76716.1; PID:g17134155; GSPDB:
A;Experimental source: strain PCC 7120
C;Genetics:
A;Gene: all5017

Query Match 63.6%; Score 35; DB 2; Length 236;
Best Local Similarity 77.8%; Pred. No. 27;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

QY 1 TRLTRKDRG 9
:|||||:
Db 94 TRNTRKDRG 102

RESULT 9
A37842
hypothetical protein 1 (xisA 3' region) - Anabaena sp. (strain PCC 7120)
C;Species: Anabaena sp.
C;Date: 21-Jun-1991 #sequence_revision 21-Jun-1991 #text_change 09-Jul-2004
C;Accession: A37842
R;Lammers, P.J.; McLaughlin, S.; Papin, S.; Trujillo-Provencio, C.; Ryncarz II, A.J.
J. Bacteriol. 172, 6981-6990, 1990
A;Title: Developmental rearrangement of cyanobacterial nif genes: nucleotide sequence,
A;Reference number: A37842; MUID:91072249; PMID:2123860
A;Accession: A37842
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-321 <LAW>
A;Cross-references: UNIPROT:P29978; GB:M38044
C;Superfamily: Anabaena hypothetical protein 1 (xisA 3' region)

Query Match 63.6%; Score 35; DB 2; Length 321;
Best Local Similarity 87.5%; Pred. No. 36;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLTRKDRG 9
:|||||:
Db 152 RLTRKDRG 159

RESULT 10
AE1987
hypothetical protein alr1448 [imported] - Nostoc sp. (strain PCC 7120)
C;Species: Nostoc sp. PCC 7120
A;Note: Nostoc sp. strain PCC 7120 is a synonym of Anabaena sp. strain PCC 7120
C;Date: 14-Dec-2001 #sequence_revision 14-Dec-2001 #text_change 09-Jul-2004
C;Accession: AE1987
R;Kaneko, T.; Nakamura, Y.; Wolk, C.P.; Kuritz, T.; Sasamoto, S.; Watanabe, A.; Iriguich
Nakazaki, N.; Shimpo, S.; Sugimoto, M.; Takazawa, M.; Yamada, M.; Yasuda, M.; Tabata, S
DNA Res. 8, 205-213, 2001
A;Title: Complete Genomic Sequence of the Filamentous Nitrogen-fixing Cyanobacterium An.
A;Reference number: AB1807; MUID:21595285; PMID:11759840
A;Accession: AE1987
A;Status: preliminary
A;Molecule type: DNA
A;Residues: 1-384 <KUR>
A;Cross-references: UNIPROT:P29978; GB:BA000019; PIDN:BAW73405.1; PID:g17130795; GSPDB:
A;Experimental source: strain PCC 7120
C;Genetics:
A;Gene: alr1448
C;Superfamily: Anabaena hypothetical protein 1 (xisA 3' region)

Query Match 63.6%; Score 35; DB 2; Length 384;
Best Local Similarity 87.5%; Pred. No. 43;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 RLTRKDRG 9
:|||||:
Db 215 RLTRKDRG 222

RESULT 11
C60950
apolipoprotein B-100 - golden hamster (fragment)
C:Species: Mesocricetus auratus (golden hamster)
C:Date: 31-Dec-1993 #sequence_revision 31-Dec-1993 #text_change 09-Jul-2004
C:Accession: C60950
R:Law, A.; Scott, J.
J. Lipid Res. 31, 1109-1120, 1990
A:Title: A cross-species comparison of the apolipoprotein B domain that binds to the LDL
A:Reference number: A60950; MUID:90324804; PMID:2373961
A:Accession: C60950
A:Molecule type: DNA
A:Residues: 1-269 <LAW>
A:Cross-references: UNIPROT:Q60537; UNIPROT:Q60536
C:Superfamily: apolipoprotein B
C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;

Query Match 62.7%; Score 34.5; DB 2; Length 269;
Best Local Similarity 81.8%; Pred. No. 38;
Matches 9; Conservative 1; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TRLTRKDRGLK 11
 :||||| |||||
Db 216 SRLTRK-RGLK 225

RESULT 12
JH0102
apolipoprotein B - golden hamster (fragment)
C:Species: Mesocricetus auratus (golden hamster)
C:Date: 17-Apr-1993 #sequence_revision 17-Apr-1993 #text_change 09-Jul-2004
C:Accession: JH0102
R:Smith, T.J.
submitted to GenBank, June 1990
A:Reference number: A38864
A:Accession: JH0102
A:Molecule type: DNA
A:Residues: 1-779 <SMI>
A:Cross-references: UNIPROT:Q60536; GB:M35187
A:Note: this is a revision to the sequence from reference JH0101
R:Smith, T.J.; Hautamaa, D.; Maeda, N.
Gene 87, 309-310, 1990
A:Title: Sequence of the putative low-density lipoprotein receptor-binding regions of ap
A:Reference number: JH0101; MUID:90236327; PMID:2332175
A:Contents: annotation
A:Note: this sequence has been revised in reference A38864
C:Genetics:
A:Gene: apoB
C:Superfamily: apolipoprotein B
C:Keywords: atherosclerosis; calcium; cholesterol metabolism; chylomicron; glycoprotein;
F:435-445/Region: receptor binding
F:646-656/Region: receptor binding

Query Match 62.7%; Score 34.5; DB 2; Length 779;
Best Local Similarity 81.8%; Pred. No. 1.1e+02;
Matches 9; Conservative 1; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TRLTRKDRGLK 11
 :||||| |||||
Db 642 SRLTRK-RGLK 651

RESULT 13
F71164
hypothetical protein PH0515 - Pyrococcus horikoshii
C:Species: Pyrococcus horikoshii
C:Date: 14-Aug-1998 #sequence_revision 14-Aug-1998 #text_change 12-Jul-2004
C:Accession: F71164
R:Kawabayashi, Y.; Sawada, M.; Horikawa, H.; Haikawa, Y.; Hino, Y.; Yamamoto, S.; Sekin
M.; Ohuku, Y.; Funahashi, T.; Tanaka, T.; Kudoh, Y.; Yamazaki, J.; Kushida, N.; Oguchi
DNA Res. 5, 55-76, 1998
A:Title: Complete sequence and gene organization of the genome of a hyper-thermophilic a

A:Reference number: A71000; MUID:98344137; PMID:9679194
A:Accession: F71164
A:Status: preliminary; nucleic acid sequence not shown; translation not shown
A:Molecule type: DNA
A:Residues: 1-195 <KAW>
A:Cross-references: UNIPROT:O58251; GB:AF000002; NID:g2236129; PIDN:BA029603.1.; PID:g37
A:Experimental source: strain OT3
A:Note: this accession replaces an interim accession for a sequence replaced by GenBank
C:Genetics:
A:Gene: PH0515

Query Match 61.8%; Score 34; DB 2; Length 195;
Best Local Similarity 60.0%; Pred. No. 35;
Matches 6; Conservative 2; Mismatches 2; Indels 0; Gaps 0;

Qy 2 RLTRKDRGLK 11
 |:|||||:
Db 135 RCSEKDRGIK 144

RESULT 14
S73019
daunorubicin resistance protein drrA - Mycobacterium leprae
N:Alternate names: L518_F2_43 protein
C:Species: Mycobacterium leprae
C:Date: 19-Mar-1997 #sequence_revision 25-Apr-1997 #text_change 16-Aug-2004
C:Accession: S73019
R:Smith, D.R.; Robison, K.
submitted to the EMBL Data Library, November 1993
A:Description: Mycobacterium leprae cosmid L518.
A:Reference number: S72591
A:Accession: S73019
A:Status: preliminary
A:Molecule type: DNA
A:Residues: 1-331 <SMI>
A:Cross-references: UNIPROT:Q49938; EMBL:U00023; NID:g467194; PIDN:AAA17362.1; PID:G9467
C:Genetics:
A:Gene: drrA
C:Superfamily: ATP-binding cassette homology
C:Keywords: ATP; nucleotide binding; P-loop
F:25-216/Domain: ATP-binding cassette homology <ABC>
F:42-49/Region: nucleotide-binding motif A (P-loop)

Query Match 61.8%; Score 34; DB 2; Length 331;
Best Local Similarity 87.5%; Pred. No. 59;
Matches 7; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 3 LTRKDRGL 10
 ||| |||||
Db 58 LTRPRDRL 65

RESULT 15
D64337
16S rRNA 5'-region hypothetical protein 1 homolog - Methanococcus jannaschii
C:Species: Methanococcus jannaschii
C:Date: 13-Sep-1996 #sequence_revision 13-Sep-1996 #text_change 09-Jul-2004
C:Accession: D64337
R:Bult, C.J.; White, O.; Olsen, G.J.; Zhou, L.; Fleischmann, R.D.; Sutton, G.G.; Blake,
; Reich, C.I.; Overbeek, R.; Kirkness, E.F.; Weinstock, K.G.; Merrick, J.M.; Glodek, A
rson, J.D.; Sadow, P.W.; Hanna, M.C.; Cotton, M.D.; Roberts, K.M.; Hurst, M.A.
Science 273, 1058-1073, 1996
A:Authors: Kaine, B.P.; Borodovsky, M.; Klenk, H.P.; Fraser, C.M.; Smith, H.O.; Woese
A:Title: Complete genome sequence of the methanogenic archaeon, Methanococcus jannaschii
A:Reference number: A64300; MUID:96337999; PMID:8688087
A:Accession: D64337
A:Status: preliminary;
A:Molecule type: DNA
A:Residues: 1-389 <BUL>
A:Cross-references: UNIPROT:Q57747; GB:U67485; GB:L77117; NID:g1591020; PIDN:AAB98286.1
C:Genetics:
A:Map position: REV282927-281758
A:Start codon: GTG

C:Superfamily: fructose-1,6-bisphosphatase, archaeal type

Query Match 61.8%; Score 34; DB 2; Length 389;
 Best Local Similarity 85.7%; Fred. No. 69;
 Matches 6; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

Qy 3 ITRKDRG 9
 :|||
 Db 380 ITRKDRG 386

Search completed: January 13, 2005, 01:52:43
 Job time : 16.8689 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: January 13, 2005, 01:15:45 ; Search time 85.8361 Seconds
(without alignments)
73.735 Million cell updates/sec

Title: US-09-823-418-19
Perfect score: 55
Sequence: 1 TRLTRKDRGLK 11

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 1825181 seqs, 575374646 residues

Total number of hits satisfying chosen parameters: 1825181

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : UniProt.02.*

1: uniprot_sprot.*

2: uniprot_trembl.*

Pred. No. is the number of results predicted by chance to have a
score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

| Result No. | Score | Query Match | Length | DB ID | Description |
|------------|-------|-------------|--------|--------------|--------------------|
| 1 | 40 | 72.7 | 250 | 2 Q8DNJ3 | Q8dnj3 streptococc |
| 2 | 40 | 72.7 | 2073 | 1 BIME EMENI | P24686 emericella |
| 3 | 39 | 70.9 | 438 | 2 Q99JK2 | Q99jk2 mus musculu |
| 4 | 39 | 70.9 | 807 | 2 Q82LJ2 | Q82lj2 salmonella |
| 5 | 38.5 | 70.0 | 414 | 2 Q7YQR5 | Q7yqr5 aocus vocif |
| 6 | 38.5 | 70.0 | 596 | 2 Q28473 | Q28473 macaca fasc |
| 7 | 38.5 | 70.0 | 3262 | 2 Q13788 | Q13788 homo sapien |
| 8 | 38.5 | 70.0 | 4563 | 1 APE HUMAN | P04114 homo sapien |
| 9 | 38.5 | 70.0 | 4563 | 2 Q7Z600 | Q7z600 homo sapien |
| 10 | 37 | 67.3 | 213 | 2 Q72PV5 | Q72pv5 leptospira |
| 11 | 37 | 67.3 | 213 | 2 Q8F6D8 | Q8f6d8 leptospira |
| 12 | 37 | 67.3 | 213 | 2 AAS70931 | Aas70931 leptospir |
| 13 | 37 | 67.3 | 838 | 2 Q800W5 | Q800w5 brachydanio |
| 14 | 37 | 67.3 | 960 | 2 Q7N7N1 | Q7n7n1 photorhabdo |
| 15 | 36 | 65.5 | 100 | 1 URE3 CORGL | Q9rhm6 corynebacte |
| 16 | 36 | 65.5 | 100 | 1 URE3 LACFE | P25931 lactobacill |
| 17 | 36 | 65.5 | 100 | 2 AAD22478 | Aad22478 lactobaci |
| 18 | 36 | 65.5 | 250 | 2 Q7N7Y0 | Q7n7y0 streptococc |
| 19 | 36 | 65.5 | 261 | 1 EMCN HUMAN | Q9ulc0 homo sapien |
| 20 | 36 | 65.5 | 283 | 2 Q7PS51 | Q7ps51 anopheles g |
| 21 | 36 | 65.5 | 340 | 2 Q8GLA1 | Q8glal streptococc |
| 22 | 36 | 65.5 | 370 | 2 Q6NDR2 | Q6ndr2 rhodopsendo |
| 23 | 36 | 65.5 | 370 | 2 CAE25486 | Ca25486 rhodopsesu |
| 24 | 36 | 65.5 | 499 | 1 GSHR ARATH | P48641 arabidopsis |
| 25 | 36 | 65.5 | 499 | 2 AAM98183 | Aam98183 arabidops |
| 26 | 36 | 65.5 | 499 | 2 AAN13086 | Aan13086 arabidops |
| 27 | 36 | 65.5 | 499 | 2 AAP68309 | Aap68309 arabidops |
| 28 | 36 | 65.5 | 682 | 2 Q6NV26 | Q6nv26 brachydanio |
| 29 | 36 | 65.5 | 682 | 2 Q6UHJ0 | Q6uhj0 brachydanio |
| 30 | 36 | 65.5 | 682 | 2 AAH68340 | Aah68340 brachydan |
| 31 | 36 | 65.5 | 1007 | 2 Q8RWG0 | Q8rwo0 arabidopsis |

| | | | | | |
|----|----|------|-----|--------------|--------------------|
| 32 | 35 | 63.6 | 78 | 2 Q9JZ39 | Q9jz39 neisseria m |
| 33 | 35 | 63.6 | 111 | 2 Q95X71 | Q95x71 caenorhabdi |
| 34 | 35 | 63.6 | 198 | 2 Q8SUL5 | Q8sul5 encephalito |
| 35 | 35 | 63.6 | 220 | 2 Q8DIV0 | Q8div0 synectococc |
| 36 | 35 | 63.6 | 236 | 2 Q8YMB7 | Q8ymb7 anabaena sp |
| 37 | 35 | 63.6 | 361 | 2 Q840G9 | Q840g9 actinobacil |
| 38 | 35 | 63.6 | 374 | 2 Q8KIF9 | Q8kip9 pseudomonas |
| 39 | 35 | 63.6 | 374 | 2 Q8KIV1 | Q8kiv1 pseudomonas |
| 40 | 35 | 63.6 | 384 | 1 Y848 ANASP | P29978 anabaena sp |
| 41 | 35 | 63.6 | 391 | 2 Q8FL8 | Q8fle8 corynebacte |
| 42 | 35 | 63.6 | 480 | 2 Q9H834 | Q9h834 homo sapien |
| 43 | 35 | 63.6 | 506 | 2 Q6Z7C0 | Q6z7c0 oryza sativ |
| 44 | 35 | 63.6 | 506 | 2 BAD07650 | Bad07650 oryza sat |
| 45 | 35 | 63.6 | 506 | 2 BAD07928 | Bad07928 oryza sat |

ALIGNMENTS

RESULT 1

| ID | Q8DNJ3 | PRELIMINARY; | PRT; | 250 AA. |
|----|--|--------------|------|---------|
| AC | Q8DNJ3; | | | |
| DT | 01-MAR-2003 (TrEMBLrel. 23, Created) | | | |
| DT | 01-OCT-2003 (TrEMBLrel. 25, Last sequence update) | | | |
| DE | ABC transporter ATP-binding protein-ferric iron transport. | | | |
| GN | Name=fecC; OrderedLocusNames=apri686; | | | |
| OS | Streptococcus pneumoniae (strain ATCC BAA-255 / R6). | | | |
| OC | Bacteria; Firmicutes; Lactobacillales; Streptococcaceae; | | | |
| OC | Streptococcus. | | | |
| OX | NCBI_TaxID=171101; | | | |
| RP | SEQUENCE FROM N.A. | | | |
| RC | STRAIN=ATCC BAA-255 / R6; | | | |
| RA | MEDLINE=21429245; PubMed=11544234; | | | |
| RA | Hoskins J., Alborn W.E. Jr., Arnold J., Blaszcak L.C., Burgett S., | | | |
| RA | DeHoff B.S., Estrem S.T., Fritz L., Fu D.-J., Fuller W., Geringer C., | | | |
| RA | Gilmour R., Glass J.S., Khoja H., Kraft A.R., Lagace R.E., | | | |
| RA | LeBlanc D.J., Lee L.N., Lefkowitz E.J., Lu J., Matsushima P., | | | |
| RA | McAhren S.M., McHenry M., McLeaster K., Mundy C.W., Niclas T.I., | | | |
| RA | Norris F.H., O'Gara M., Peery R.B., Robertson G.T., Rockey P., | | | |
| RA | Sun P.-M., Winkler M.E., Yang Y., Young-Bellido M., Zhao G., | | | |
| RA | Zook C.A., Baltz R.H., Jaskunas S.R., Rostek P.R. Jr., Skatrud P.L., | | | |
| RA | Glass J.I.; | | | |
| RL | "Genome of the bacterium Streptococcus pneumoniae strain R6."; | | | |
| CC | J. Bacteriol. 183:5709-5717(2001). | | | |
| CC | -I- SIMILARITY: Belongs to the ABC transporter family. | | | |
| DR | EMBL; AE008534; AAL00489.1; -. | | | |
| DR | PIR; D98082; D98082. | | | |
| DR | HSSP; P06611; 1L7V. | | | |
| DR | GO; GO:0016020; C:membrane; IEA. | | | |
| DR | GO; GO:0005534; F:ATP binding; IEA. | | | |
| DR | GO; GO:0004009; F:ATP-binding cassette (ABC) transporter acti. .; IEA. | | | |
| DR | GO; GO:0000166; F:nucleotide binding; IEA. | | | |
| DR | GO; GO:0006810; P:transport; IEA. | | | |
| DR | InterPro; IPR003593; AAA_ATPase. | | | |
| DR | InterPro; IPR003439; ABC_transporter. | | | |
| DR | Pfam; PF00005; ABC_tran; 1. | | | |
| DR | PRODOM; PD000006; ABC_transporter; 1. | | | |
| DR | SMART; SM00382; AAA; 1. | | | |
| DR | PROSITE; PS00211; ABC_TRANSPORTER_1; 1. | | | |
| DR | PROSITE; PS00893; ABC_TRANSPORTER_2; 1. | | | |
| KW | ATP-binding; Complete proteome. | | | |
| SQ | SEQUENCE 250 AA; 28528 MW; FFC841A29F30033D CRC64; | | | |

| | | | | |
|-----------------------|--------|----------------|-------|---------------|
| Query Match | 72.7%; | Score 40; | DB 2; | Length 250; |
| Best Local Similarity | 70.0%; | Pred. No. 7.2; | | |
| Matches | 7; | Conservative | 3; | Mismatches 0; |
| | | | | Indels 0; |
| | | | | Gaps 0; |
| Qy | 1 | TRLTRKDRGL | 10 | |
| | | | | |
| Db | 47 | SLRTRKDRGV | 56 | |

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RESULT 2
BIME EMENI
ID BIME EMENI STANDARD; PRT; 2073 AA.
AC F24686;
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 01-MAR-1992 (Rel. 21, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Negative regulator of mitosis.
GN Name=BIME;
OS Emericella nidulans (Aspergillus nidulans).
OC Eukaryota; Fungi; Ascomycota; Pezizomycotina; Eurotiomycetes;
OC Eurotiales; Trichocomaceae; Emericella.
OX NCBI_TaxID=162425;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90375468; PubMed=1697851;
RA Engle D.B.; Osmani S.A.; Osmani A.H.; Rosborough S.; Xiang X.;
RA Morris N.R.;
RT "A negative regulator of mitosis in Aspergillus is a putative
RT membrane-spanning protein.";
RL J. Biol. Chem. 265:16132-16137(1990).
CC -!- FUNCTION: Negative regulator of mitosis in E.nidulans. This
CC protein is part of a regulatory pathway that includes the nimA
CC protein kinase. It is required to prevent premature entry into
CC mitosis. Mutations to this protein both cause cells to enter
CC mitosis and prevent them from leaving mitosis.
CC -!- SIMILARITY: Contains 4 PC repeats.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC entities requires a license agreement (see http://www.isb-sib.ch/announce/
CC or send an email to license@isb-sib.ch).
CC -----
DR EMBL; M59705; AAA51478.1; -.
DR PIR; A37879; BWASBE.
DR InterPro; IPR002015; APC_proteasome.
DR Pfam; PF01851; PC_rep; 4.
KW Mitosis; Repeat; Transmembrane.
FT DOMAIN 342 353 Nuclear localization signal (Potential).
FT TRANSMEM 1623 1643 Potential.
FT TRANSMEM 1685 1703 Potential.
FT TRANSMEM 1746 1764 Potential.
SQ SEQUENCE 2073 AA; 229178 MW; 05E4E81EADDF51E4 CRC64;

Query Match 72.7%; Score 40; DB 1; Length 2073;
Best Local Similarity 80.0%; Pred. No. 88;
Matches 8; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy 1 TRLTRKDRGL 10
|||||: |||
Db 832 TRLTRKDRGL 841

RESULT 3
Q99JK2 PRELIMINARY; PRT; 438 AA.
AC Q99JK2;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Lars protein.
GN Name=Lars;
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.

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RC STRAIN=Czech II;
RC TISSUE=Mammary tumor metastasized to lung. Tumor arose spontaneously;
RX MEDLINE=22388257; PubMed=12477932;
RA Strausberg R.L.; Feingold E.A.; Grouse L.H.; Derge J.G.;
RA Klausner R.D.; Collins F.S.; Wagner L.; Shenmen C.M.; Schuler G.D.;
RA Altschul S.F.; Zeeberg B.; Buetow K.H.; Schaefer C.F.; Bhat N.K.;
RA Hopkins R.F.; Jordan H.; Moore T.; Max S.I.; Wang J.; Haieh P.;
RA Diatchenko L.; Marusina K.; Farmer A.A.; Rubin G.M.; Hong L.;
RA Stapleton M.; Soares M.B.; Bonaldo M.F.; Casavant T.L.; Scheetz T.E.;
RA Brownstein M.J.; Uesdin T.B.; Toshiyuki S.; Carninci P.; Prange C.;
RA Raha S.S.; Loquellano N.A.; Peters G.J.; Abramson R.D.; Mullaly S.J.;
RA Bosak S.A.; McEwan P.J.; McKernan K.J.; Malek J.A.; Gunaratne P.H.;
RA Richards S.; Worley K.C.; Hale S.; Garcia A.M.; Gay L.J.; Hulyk S.W.;
RA Villalon D.K.; Muzny D.W.; Sodergren E.J.; Lu X.; Gibbs R.A.;
RA Fahney J.; Helton E.; Kettelman M.; Madan A.; Rodrigues S.; Sanchez A.;
RA Whiting M.; Madan A.; Young A.C.; Shevchenko Y.; Bouffard G.G.;
RA Blakesley R.C.; Touchman J.W.; Green E.D.; Dickson M.C.;
RA Rodriguez A.C.; Grimwood J.; Schmitz J.; Myers R.M.; Butterfield V.S.;
RA Krzywinski M.I.; Skaleka U.; Smailus D.E.; Schnerch A.; Schein J.E.;
RA Jones S.J.; Marra M.A.;
RT "Generation and initial analysis of more than 15,000 full-length human
RT and mouse cDNA sequences.";
RL Proc. Natl. Acad. Sci. U.S.A. 99:16899-16903(2002).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=Czech II;
RC TISSUE=Mammary tumor metastasized to lung. Tumor arose spontaneously;
RA Strausberg R.;
RL EMBL; BC006060; AAH06060.1; -.
DR MGD; MGI:1913808; Lars.
DR InterPro; IPR002088; PPTA.
DR InterPro; IPR009080; TRNAsyn_la_bind.
DR PROSITE; PS00904; PPTA; UNKNOWN 1.
SQ SEQUENCE 438 AA; 49840 MW; 2E730A99D65AFBF0 CRC64;

Query Match 70.9%; Score 39; DB 2; Length 438;
Best Local Similarity 70.0%; Pred. No. 23;
Matches 7; Conservative 2; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLTRKDRGLK 11
|||||: |||
Db 361 RLTKDRGK 370

RESULT 4
Q8Z1J2 PRELIMINARY; PRT; 807 AA.
AC Q8Z1J2; Q7C5G1;
DT 01-MAR-2002 (TrEMBLrel. 20, Created)
DT 01-MAR-2002 (TrEMBLrel. 20, Last sequence update)
DT 01-OCT-2004 (TrEMBLrel. 28, Last annotation update)
DE Hypothetical protein STY4573.
DE OrderedLocusNames=STY4573, t4270;
OS Salmonella typhi.
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Salmonella.
OX NCBI_TaxID=601;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21534947; PubMed=11677608; DOI=10.1038/35101607;
RA Parkhill J.; Dougan G.; James K.D.; Thomson N.R.; Pickard D.; Wain J.;
RA Churcher C.M.; Mungall K.L.; Bentley S.D.; Holden M.T.G.; Sebahia M.;
RA Baker S.; Basham D.; Brooks K.; Chillingworth T.; Connor P.;
RA Cronin A.; Davis P.; Davies R.M.; Dowd L.; White N.; Farrar J.;
RA Feltwell T.; Hamlin N.; Haque A.; Hien T.T.; Holroyd S.; Jagels K.;
RA Krogh A.; Larsen T.S.; Leather S.; Moule S.; O'Gaora P.; Parry C.;
RA Quail M.A.; Rutherford K.M.; Simmonds M.; Skelton J.; Stevens K.;
RA Whitehead S.; Barrell B.G.;
RT "Complete genome sequence of a multiple drug resistant Salmonella
RT enterica serovar Typhi CT18.";
RL Nature 413:848-852(2001).

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RN SEQUENCE FROM N.A.
RP STRAIN=Ty2 / ATCC 700931;
RX MEDLINE=22531367; PubMed=12644504;
RA Deng W., Liou S.-R., Plunkett G. III, Mayhew G.F., Rose D.J.,
RA Burland V., Kodoyianni V., Schwartz D.C., Blattner F.R.;
RT "Comparative genomics of Salmonella enterica serovar Typhi strains Ty2
RT and CT18.";
RL J. Bacteriol. 185:2330-2337(2003).
DR EMBL; AL627282; CAD09348.1; -.
DR EMBL; AE016848; AA071728.1; -.
KW Complete proteome; Hypothetical protein.
SQ SEQUENCE 807 AA; 90898 MW; 577CIDA31611BCF0 CRC64;

Query Match 70.9%; Score 39; DB 2; Length 807;
Best Local Similarity 88.9%; Pred. No. 47;
Matches 8; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 2 RLTRKDRGL 10
Db 508 RLTRADRGL 516
||||| |||||

RESULT 5
Q7YQRS PRELIMINARY; PRT; 414 AA.
AC Q7YQRS;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-OCT-2003 (TrEMBLrel. 25, Last annotation update)
DE Apolipoprotein B 100 (Fragment).
GN Name=apob-100;
OS Aotus vociferans (Spix's owl monkey).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Platyrrhini; Cebidae; Aotinae; Aotus.
OX NCBI_TaxID=57176;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=22761261; PubMed=12878460;
RA Amrine-Madsen H., Koepfli K.-P., Wayne R.K., Springer M.S.;
RT "A new phylogenetic marker, apolipoprotein B, provides compelling
RT evidence for eutherian relationships.";
RL Mol. Phylogenet. Evol. 28:225-240(2003).
DR EMBL; AF548396; AAP97352.1; -.
KW Lipoprotein.
FT NON TER 1 1
FT NON TER 414 414
SQ SEQUENCE 414 AA; 45955 MW; EFA8492157E1BDE CRC64;

Query Match 70.0%; Score 38.5; DB 2; Length 414;
Best Local Similarity 90.9%; Pred. No. 27;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TLTRKDRGLK 11
Db 258 TLTRK-RGLK 267
||||| |||||

RESULT 6
Q28473 PRELIMINARY; PRT; 596 AA.
AC Q28473;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE Apolipoprotein B (Fragment).
OS Macaca fascicularis (Crab eating macaque) (Cynomolgus monkey).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Cercopithecoidea;
OC Cercopithecinae; Macaca.
OX NCBI_TaxID=9541;
RN [1]
RP SEQUENCE FROM N.A.

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RC TISSUE=Liver;
RX MEDLINE=92075708; PubMed=1742325;
RA Pape M.E., Castle C.K., Murray R.W., Funk G.M., Hunt C.E.,
RA Marotti K.R., Melchior G.W.;
RT "Apo B metabolism in the cynomolgus monkey: evidence for post-
RT transcriptional regulation.";
RT Biochim. Biophys. Acta 1086:326-334(1991).
RN [2]
RL SEQUENCE FROM N.A.
RP TISSUE=Liver;
RC TISSUE=Liver;
RA Murray R.;
RL Submitted (PEB-1992) to the EMBL/GenBank/DBSJ databases.
DR EMBL; X15737; CAA33755.1; -.
DR PIR; S32802; S32802.
KW Lipoprotein.
FT NON TER 1 1
FT NON TER 596 596
SQ SEQUENCE 596 AA; 66757 MW; B13BBA74E25C3120 CRC64;

Query Match 70.0%; Score 38.5; DB 2; Length 596;
Best Local Similarity 90.9%; Pred. No. 42;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TLTRKDRGLK 11
Db 226 TLTRK-RGLK 235
||||| |||||

RESULT 7
Q13788 PRELIMINARY; PRT; 3262 AA.
AC Q13788;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE APOB protein (Fragment).
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=8719199; PubMed=2883086;
RA Carlsson P., Darnfors C., Olsson S.O., Bjursell G.;
RT "Analysis of the human apolipoprotein B gene; complete structure of
RT the B-74 region.";
RL Gene 49:29-51(1986).
DR EMBL; M15421; AAA51758.1; -.
DR PIR; A27850; LPHUB.
DR GO; GO:0005576; C:extracellular; NAS.
DR GO; GO:0005319; F:lipid transporter activity; NAS.
DR GO; GO:0006869; P:lipid transport; NAS.
FT NON TER 1 1
SQ SEQUENCE 3262 AA; 370140 MW; 56603BC0618DD40D CRC64;

Query Match 70.0%; Score 38.5; DB 2; Length 3262;
Best Local Similarity 90.9%; Pred. No. 3.2e+02;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TLTRKDRGLK 11
Db 2084 TLTRK-RGLK 2093
||||| |||||

RESULT 8
APB HUMAN
ID APB HUMAN STANDARD; PRT; 4563 AA.
AC P04114; O00502; Q13787;
DT 01-NOV-1986 (Rel. 03, Created)
DT 01-NOV-1986 (Rel. 03, Last sequence update)
DT 05-JUL-2004 (Rel. 44, Last annotation update)
DE Apolipoprotein B-100 precursor (Apo B-100) (Contains: Apolipoprotein

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DE B-48 (Apo B-48)].
GN Name=APOB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=87016385; PubMed=3763409;
RA Knott T.C., Wallis S.C., Powell L.M., Pease R.J., Lueis A.J.,
RA Blackhart B., McCarthy B.J., Mahley R.W., Levy-Wilson B., Scott J.;
RT "Complete cDNA and derived protein sequence of human apolipoprotein B-
RT 100.";
RL Nucleic Acids Res. 14:7501-7503(1986).
RN [2]
RP SEQUENCE FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=98003974; PubMed=3652907;
RA Ludwig E.H., Blackhart B.D., Pierotti V.R., Caiati L., Fortier C.,
RA Knott T., Scott J., Mahley R.W., Levy-Wilson B., McCarthy B.J.;
RT "DNA sequence of the human apolipoprotein B gene.";
RL Nucleic Acids Res. 14:3663-372(1987).
RN [3]
RP SEQUENCE FROM N.A., AND VARIANTS ILE-98 AND GLU-4181.
RX MEDLINE=87008488; PubMed=3759943;
RA Chen S.-H., Yang C.-Y., Chen P.-F., Setzer D., Tanimura M., Li W.-H.,
RA Gotto A.M. Jr., Chan L.;
RT "The complete cDNA and amino acid sequence of human apolipoprotein B-
RT 100.";
RL J. Biol. Chem. 261:12918-12921(1986).
RN [4]
RP SEQUENCE FROM N.A.
RX MEDLINE=87041416; PubMed=3464946;
RA Law S.W., Grant S.M., Higuchi K., Hospattankar A.V., Lackner K.J.,
RA Lee N., Brewer H.B. Jr.;
RT "Human liver apolipoprotein B-100 cDNA: complete nucleic acid and
RT derived amino acid sequence.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:8142-8146(1986).
RN [5]
RP SEQUENCE FROM N.A.
RX MEDLINE=87161758; PubMed=3030729;
RA Cladaras C., Hadzopoulou-Cladaras M., Nolte R.T., Atkinson D.,
RA Zannis V.I.;
RT "The complete sequence and structural analysis of human apolipoprotein
RT B-100: relationship between apoB-100 and apoB-48 forms.";
RL EMBO J. 5:3495-3507(1986).
RN [6]
RP SEQUENCE OF 709-906 FROM N.A.
RX MEDLINE=85270450; PubMed=3860836;
RA Deeb S.S., Motulsky A.G., Albers J.J.;
RT "A partial cDNA clone for human apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 82:4983-4986(1985).
RN [7]
RP SEQUENCE OF 3056-3159 FROM N.A.
RX MEDLINE=86041888; PubMed=3903660;
RA Mehrabian M., Schumaker V.N., Fareed G.C., West R., Johnson D.F.,
RA Kirchgesner T.G., Lin H.-C., Wang X., Ma Y., Mendiaz E., Lusis A.J.;
RT "Human apolipoprotein B: identification of cDNA clones and
RT characterization of mRNA.";
RL Nucleic Acids Res. 13:6937-6953(1985).
RN [8]
RP SEQUENCE OF 1937-2018 AND 3811-4334 FROM N.A., AND VARIANT GLU-4181.
RX MEDLINE=86093680; PubMed=3841204;
RA Carlsson P., Olofsson S.O., Bondjers G., Darnfors C., Wiklund O.,
RA Bjursell G.;
RT "Molecular cloning of human apolipoprotein B cDNA.";
RL Nucleic Acids Res. 13:8813-8826(1985).
RN [9]
RP SEQUENCE OF 3109-4563 FROM N.A.
RX MEDLINE=85300528; PubMed=2994222;
RA Knott T.J., Rall S.C. Jr., Innerarity T.L., Jacobson S.F., Urdea M.S.,
RA Levy-Wilson B., Powell L.M., Pease R.J., Eddy R., Nakai H., Byers M.,
RA Priestley L.M., Robertson E., Rall L.B., Betscholtz C., Shows T.B.,
RA Mahley R.W., Scott J.;

RT "Human apolipoprotein B: structure of carboxyl-terminal domains, sites
RT of gene expression, and chromosomal localization.";
RN Science 230:37-43(1985).
RN [10]
RP SEQUENCE OF 1-291 FROM N.A.
RX MEDLINE=86149325; PubMed=3513177;
RA Protter A.A., Hardman D.A., Schilling J.W., Miller J., Appleby V.,
RA Chen G.C., Kirsher S.W., McEnroe G., Kane J.P.;
RT "Isolation of a cDNA clone encoding the amino-terminal region of human
RT apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:1467-1471(1986).
RN [11]
RP SEQUENCE OF 1-1670 FROM N.A., AND VARIANT ILE-98.
RX MEDLINE=86287319; PubMed=3461454;
RA Protter A.A., Hardman D.A., Sato K.Y., Schilling J.W., Yamanaka M.,
RA Hort Y.J., Hjerrild K.A., Chen G.C., Kane J.P.;
RT "Analysis of cDNA clones encoding the entire B-26 region of human
RT apolipoprotein B.";
RL Proc. Natl. Acad. Sci. U.S.A. 83:5678-5682(1986).
RN [12]
RP PARTIAL SEQUENCE, AND IDENTIFICATION OF APO-B48.
RX MEDLINE=88018019; PubMed=3659919;
RA Chen S.-H., Habib G., Yang C.-H., Gu Z.-W., Lee B.R., Weng S.-H.,
RA Silberman S.R., Cai S.-J., Deslypere J.P., Rosseneu M.,
RA Gotto A.M. Jr., Li W.-H., Chan L.;
RT "Apolipoprotein B-48 is the product of a messenger RNA with an organ-
RT specific in-frame stop codon.";
RL Science 238:363-366(1987).
RN [13]
RP DOMAINS.
RX MEDLINE=87039351; PubMed=3773997;
RA Knott T.C., Pease R.J., Powell L.M., Wallis S.C., Rall S.C. Jr.,
RA Innerarity T.L., Blackhart B., Taylor W.R., Marcel Y., Milne R.,
RA Johnson D., Fuller M., Lusis A.J., McCarthy B.J., Mahley R.W.,
RA Levy-Wilson B., Scott J.;
RT "Complete protein sequence and identification of structural domains of
RT human apolipoprotein B.";
RL Nature 323:734-738(1986).
RN [14]
RP DOMAINS.
RX Yang C.-Y., Chen S.-H., Gianturco S.H., Bradley W.A., Sparrow J.T.,
RA Tanimura M., Li W.-H., Sparrow D.A., Deloof H., Rosseneu M.,
RA Lee F.-S., Gu Z.-W., Gotto A.M. Jr., Chan L.;
RT "Sequence, structure, receptor-binding domains and internal repeats of
RT human apolipoprotein B-100.";
RL Nature 323:738-742(1986).
RN [15]
RP CALCIUM-BINDING DATA.
RX MEDLINE=86242245; PubMed=3087360;
RA Dashti N., Lee D.M., Mok T.;
RT "Apolipoprotein B is a calcium binding protein.";
RL Biochem. Biophys. Res. Commun. 137:493-499(1986).
RN [16]
RP PALMITOYLATION OF CVS-1112.
RX MEDLINE=20143590; PubMed=10679026;
RA Zhao Y., McCabe J.B., Vance J., Berthiaume L.G.;
RT "Palmitoylation of apolipoprotein B is required for proper
RT intracellular sorting and transport of cholesterol esters and
RT triglycerides.";
RL Mol. Biol. Cell 11:721-734(2000).
RN [17]
RP VARIANT SER-4338.
RX MEDLINE=91071750; PubMed=1979313;
RA Navajas M., Laurent A.-M., Moreel J.-F., Ragab A., Cambou J.-P.,
RA Cuny G., Cambien F., Roizes G.;
RT "Detection by denaturing gradient gel electrophoresis of a new
RT polymorphism in the apolipoprotein B gene.";
RL Hum. Genet. 86:91-93(1990).
RN [18]
RP VARIANT FDS GLN-3527.
RX MEDLINE=89098975; PubMed=2563166;
RA Soria L.F., Ludwig E.H., Clarke H.R.G., Vega G.L., Grundy S.M.,
RA McCarthy B.J.;

RT "Association between a specific apolipoprotein B mutation and familial
RT defective apolipoprotein B-100.";
RL Proc. Natl. Acad. Sci. U.S.A. 86:587-591(1989).
RN [19]
RP VARIANT LEU-2739.
RX MEDLINE=91016974; PubMed=2216805;
RA Huang L.-S., Gavish D., Breslow J.L.;
RT "Sequence polymorphism in the human apoB gene at position 8344.";
RL Nucleic Acids Res. 18:5922-5922(1990).
RN [20]
RP VARIANT FDB CYS-3558.
RX MEDLINE=95190020; PubMed=7883971;
RA Pullinger C.R., Hennessy L.K., Chatterton J.E., Liu W., Love J.A.,
Mendel C.M., Frost P.H., Malloy M.J., Schumaker V.N., Kane J.P.;
RT "Familial ligand-defective apolipoprotein B. Identification of a new
RT mutation that decreases LDL receptor binding affinity.";
RL J. Clin. Invest. 95:1225-1234(1995).
RN [21]
RP VARIANTS LEU-1437; SER-1914; LYS-2566; THR-3121; ALA-3945; MET-4128
AND THR-4481.
RX MEDLINE=97044521; PubMed=8889592;
RA Poirier O., Ricard S., Behague I., Souriau C., Evans A.E.,
Arveiler D., Marques-Vidal P., Luc G., Roizes G., Cambien F.;
RT "Detection of new variants in the apolipoprotein B (Apo B) gene by
RT PCR-SSCP.";
RL Hum. Mutat. 8:282-285(1996).
RN [22]
RP VARIANTS FDB GLN-3527 AND CYS-3558.
RX MEDLINE=97403938; PubMed=9259199;
RA Rabes J.P., Varret M., Saint-Jore B., Erlich D., Jondeau G.,
Kremf M., Giraudet P., Junien C., Boileau C.;
RT "Familial ligand-defective apolipoprotein B-100: simultaneous
RT detection of the ARG3500-->GLN and ARG3531-->CYS mutations in a French
RT population.";
RL Hum. Mutat. 10:160-163(1997).
RN [23]
RP VARIANTS SER-1914; ARG-1923; LEU-2739; ASP-3319; THR-3427; GLN-3432
AND ILE-3921.
RX MEDLINE=98141125; PubMed=9490296;
RA Leren T.P., Bakken K.S., Hoel V., Hjermann I., Berg K.;
RT "Screening for mutations of the apolipoprotein B gene causing
RT hypocholesterolemia.";
RL Hum. Genet. 102:44-49(1998).
CC -!- FUNCTION: Apolipoprotein B is a major protein constituent of
CC chylomicrons (apo B-48), LDL (apo B-100) and VLDL (apo B-100). Apo
CC B-100 functions as a recognition signal for the cellular binding
CC and internalization of LDL particles by the apoB/E receptor.
CC -!- SUBCELLULAR LOCATION: Secreted.

Query Match 70.0%; Score 38.5; DB 1; Length 4563;
Best Local Similarity 90.9%; Pred. No. 4.7e+02;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TRLTRKDRGLK 11
Db 3385 TRLTRK-RGLK 3394

RESULT 9
Q7Z600 PRELIMINARY; PRT; 4563 AA.
AC Q7Z600;
DT 01-OCT-2003 (TrEMBLrel. 25, Created)
DT 01-OCT-2003 (TrEMBLrel. 25, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Apolipoprotein B (Including Ag(X) antigen).
GN Name=ApoB;
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.

RA Rieder M.J., Carrington D.P., da Ponte S.H., Hastings N.C.,
RA Ahearn M.O., Kuldane S.A., Rajkumar N., Toth E.J., Yi Q.,
RA Nickerson D.A.;
RL Submitted (JUN-2003) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY324608; AAP72970.1; -;
DR GO; GO:0005319; F:lipid transporter activity; IEA.
DR GO; GO:0006869; P:lipid transport; IEA.
DR InterPro; IPR009454; DUF1081.
DR InterPro; IPR001747; Lipid_transprt_N.
DR Pfam; PF06448; DUF1081; 1;
DR SMART; SM00638; LPD_N; 1.
DR Pfam; PF01347; Vitellogenin_N; 1.
DR Lipoprotein.
KW Lipoprotein.
SQ SEQUENCE 4563 AA; 515553 MW; 030B34167CEDC63C CRC64;

Query Match 70.0%; Score 38.5; DB 2; Length 4563;
Best Local Similarity 90.9%; Pred. No. 4.7e+02;
Matches 10; Conservative 0; Mismatches 0; Indels 1; Gaps 1;

Qy 1 TRLTRKDRGLK 11
Db 3385 TRLTRK-RGLK 3394

RESULT 10
Q7ZPV5 PRELIMINARY; PRT; 213 AA.
ID Q7ZPV5
AC Q7ZPV5;
DT 05-JUL-2004 (TrEMBLrel. 27, Created)
DT 05-JUL-2004 (TrEMBLrel. 27, Last sequence update)
DT 05-JUL-2004 (TrEMBLrel. 27, Last annotation update)
DE 3-methyladenine DNA glycosylase.
GN Name=alka; OrderedLocNames=LIC12362;
OS Leptospira interrogans (serogroup Icterohaemorrhagiae / serovar
OS Copenhagen).
OC Bacteria; Spirochaetes; Spirochaetales; Leptospiraceae; Leptospira.
OX NCBI_TaxID=44275;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Fiocruz LI-130;
RX PubMed=15028702;
RA Nascimento A.L.T.O., Ko A.I., Martins E.A.L., Monteiro-Vitorello C.B.,
Ho P.L., Haake D.A., Verjovski-Almeida S., Hartskeerl R.A.,
Marques M.V., Oliveira M.C., Menck C.F.M., Leite L.C.C., Carrer H.,
Coutinho L.L., Degraeve W.M., Dellagostin O.A., El-Dorry H.,
Ferre E.S., Ferro M.I.T., Furian L.R., Gamberini M., Giglioti E.A.,
Goes-Neto A., Goldman G.H., Goldman M.H.S., Harakava R.,
Jeronimo S.M.B., Junqueira-de-Azevedo I.L.M., Kimura E.T.,
Kuramae E.E., Lemos E.G.M., Lemos M.V.F., Marino C.L., Nunes L.R.,
de Oliveira R.C., Pereira G.G., Reis M.S., Schriefer A.,
Siqueira W.J., Sommer P., Tsai S.M., Simpson A.J.G., Ferro J.A.,
Camargo L.E.A., Kitajima J.P., Setubal J.C., Van Sluys M.A.;
RT "Comparative genomics of two Leptospira interrogans serovars reveals
RT novel insights into physiology and pathogenesis.";
RL J. Bacteriol. 186:2164-2172(2004).
DR EMBL; AE017296; AAS70931.1; -;
DR InterPro; IPR011257; DNA_glycylase.
DR InterPro; IPR003265; Endo_3c.
DR Pfam; PF00730; Hnh-GPD; 1.
DR SMART; SM00478; ENDO3c; 1.
KW Complete proteome.
SQ SEQUENCE 213 AA; 24639 MW; 73E6CA3B0C737487 CRC64;

Query Match 67.3%; Score 37; DB 2; Length 213;
Best Local Similarity 77.8%; Pred. No. 26;
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;

Qy 2 RLTRKDRGL 10
Db 19 QLSRKDRGL 27

RESULT 11

Q8F6D8
ID Q8F6D8 PRELIMINARY; PRT; 213 AA.
AC Q8F6D8
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-JUN-2003 (TrEMBLrel. 24, Last annotation update)
DE DNA-3-methyladenine glycosylase (EC 3.2.2.21).
GN Name=ag1; OrderedLocusNames=LAI1370;
OS Leptospira interrogans.
OC Bacteria; Spirochaetes; Spirochaetales; Leptospiraceae; Leptospira.
OX NCBI_TaxID=173;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=56601 / Serogroup Icterohaemorrhagiae / Serovar lai;
RX MEDLINE=22598143; PubMed=12712204;
RA Ren S.-X., Fu G., Jiang X.-G., Zeng R., Miao Y.-G., Xu H.,
RA Zhang Y.-X., Xiong H., Lu G., Lu L.-F., Jiang H.-Q., Jia J., Tu Y.-F.,
RA Jiang J.-X., Gu W.-Y., Zhang Y.-Q., Cai Z., Sheng H.-H., Yin H.-F.,
RA Zhang Y., Zhu G.-F., Wan M., Huang H.-L., Qian Z., Wang S.-Y., Ma W.,
RA Yao Z.-J., Shen Y., Qiang B.-Q., Xia Q.-C., Guo X.-K., Danchin A.,
RA Saint Girons I., Somerville R.L., Wen Y.-W., Shi M.-H., Chen Z.,
RA Xu J.-G., Zhao G.-P.;
RT "Unique physiological and pathogenic features of Leptospira
interrogans revealed by whole-genome sequencing.";
RL Nature 422:888-893(2003).
DR ENBL, AE011316; ANA8569.1; -
DR GO: GO:0003905; F:alkylase DNA N-glycosylase activity; IEA.
DR GO: GO:0016798; F:hydrolase activity, acting on glycosyl bonds; IEA.
DR GO: GO:0006284; P:base-excision repair; IEA.
DR GO: GO:0005975; P:carbohydrate metabolism; IEA.
DR InterPro: IPR011257; DNA_glycosylase.
DR InterPro: IPR003265; Endo_3c.
DR Pfam: PF00730; HNH-GPD; 1.
DR SMART: SM00478; ENDO3c; 1.
DR Complete proteome; Glycosidase; Hydrolase.
KW SEQUENCE 213 AA; 24567 MW; 01C4CA3DB9259732 CRC64;
Query Match 67.3%; Score 37; DB 2; Length 213;
Best Local Similarity 77.8%; Pred. No. 26;
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 2 RLTRKDRGL 10
Db 19 QLSRKDRGL 27

RESULT 12
AAS70931 PRELIMINARY; PRT; 213 AA.
AC AAS70931
DT 24-MAR-2004 (TrEMBLrel. 27, Created)
DT 24-MAR-2004 (TrEMBLrel. 27, Last sequence update)
DT 11-MAY-2004 (TrEMBLrel. 27, Last annotation update)
DE 3-methyladenine DNA glycosylase.
GN ALKA OR LIC12362.
OS Leptospira interrogans (serogroup Icterohaemorrhagiae / serovar
Copenhagen).
OC Bacteria; Spirochaetes; Spirochaetales; Leptospiraceae; Leptospira.
OX NCBI_TaxID=44275;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=F10ctruz L1-130;
RX PubMed=15028702;
RA Nascimento A.L.F.O., Ko A.I., Martins E.A.L., Monteiro-Vitorello C.B.,
RA Ho P.L., Haake D.A., Verjovski-Almeida S., Hartskeerl R.A.,
RA Marques M.V., Oliveira M.C., Mencia C.F.M., Leite L.C.C., Carrer H.,
RA Coutinho L.L., Degreve W.M., Dellagostin O.A., El-Dorri H.,
RA Ferro E.S., Ferro M.I.T., Furlan L.R., Gamberini M., Gigliotti E.A.,
RA Goes-Neto A., Goldman G.H., Goldman M.H.S., Harakava R.,
RA Jeronimo S.M.B., Junqueira-de-Azevedo J.L.M., Kimura E.T.,
RA Kuramae E.E., Lemos E.G.M., Lemos M.V.F., Marino C.L., Nunes L.R.,
RA de Oliveira R.C., Pereira G.G., Reis M.S., Schriefer A.,
RA Siqueira W.J., Sommer P., Tsai S.M., Simpson A.J.G., Ferro J.A.,

RA Camargo L.E.A., Kitajima J.P., Setubal J.C., Van Sluys M.A.;
RT "Comparative genomics of two Leptospira interrogans serovars reveals
novel insights into physiology and pathogenesis.";
RL J. Bacteriol. 186:2164-2172(2004).
DR ENBL, AE017296; AAS70931.1; -
SQ SEQUENCE 213 AA; 24639 MW; 73B6CA3E0C737487 CRC64;
Query Match 67.3%; Score 37; DB 2; Length 213;
Best Local Similarity 77.8%; Pred. No. 26;
Matches 7; Conservative 2; Mismatches 0; Indels 0; Gaps 0;
QY 2 RLTRKDRGL 10
Db 19 QLSRKDRGL 27

RESULT 13
Q800W5 PRELIMINARY; PRT; 838 AA.
ID Q800W5
AC Q800W5
DT 01-JUN-2003 (TrEMBLrel. 24, Created)
DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Frodo2.
GN Name=frd2;
OS Brachydanio rerio (Zebrafish) (Danio rerio).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Actinopterygii; Neopterygii; Teleostei; Ostariophysi; Cypriniformes;
OC Cyprinidae; Danio.
OX NCBI_TaxID=7955;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=21984492; PubMed=11941372;
RA Gloy J., Hikasa H., Sokol S.Y.;
RT "Frodo interacts with Dishevelled to transduce Wnt signals.";
RL Nat. Cell Biol. 4:351-357(2002).
RN [2]
RP SEQUENCE FROM N.A.
RX PubMed=15188426;
RA Gillhouse M., Wagner Nyholm M., Hikasa H., Sokol S.Y., Grinblatt Y.;
RT "Two Frodo/Dapper homologs are expressed in the developing brain and
mesoderm of zebrafish.";
RL Dev. Dyn. 230:403-409(2004).
RN [3]
RP SEQUENCE FROM N.A.
RA Hikasa H., Sokol S.Y.;
RL Submitted (JAN-2004) to the EMBL/GenBank/DBJ databases.
DR ENBL; AY208969; AAO49711.2; -
DR ZFIN; ZDB-GENE-030131-9975; dact.
SQ SEQUENCE 838 AA; 91349 MW; 6906B5ECC35012BC CRC64;
Query Match 67.3%; Score 37; DB 2; Length 838;
Best Local Similarity 72.7%; Pred. No. 1.3e+02;
Matches 8; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 1 TLTRKDRGLK 11
Db 86 TLTRKDRGLK 96

RESULT 14
Q7N7N1 PRELIMINARY; PRT; 960 AA.
ID Q7N7N1
AC Q7N7N1
DT 01-MAR-2004 (TrEMBLrel. 26, Created)
DT 01-MAR-2004 (TrEMBLrel. 26, Last sequence update)
DT 01-MAR-2004 (TrEMBLrel. 26, Last annotation update)
DE Similar to unknown protein.
GN OrderedLocusNames=plu1085;
OS Photobacterium luminescens (subsp. laumondii).
OC Bacteria; Proteobacteria; Gammaproteobacteria; Enterobacteriales;
OC Enterobacteriaceae; Photobacterium.
OX NCBI_TaxID=141679;


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RN  SEQUENCE FROM N.A.
RC  STRAIN=TT01;
RX  MEDLINE=22957627; PubMed=14528314;
RA  Duchaud E., Rusniok C., Frangeul L., Buchrieser C., Givaudan A.,
RA  Taourit S., Bocs S., Boursaux-Eude C., Chandler M., Charles J.-F.,
RA  Daasa E., Deroose R., Derzelle S., Freysinet G., Gaudrault S.,
RA  Medigue M., Lanois A., Powell K., Sigulier P., Vincent R., Wingate V.,
RA  Zouine M., Glaser P., Boemare N., Danchin A., Kunat F.;
RT  "The genome sequence of the entomopathogenic bacterium Photobacterium
RT  luminescens.";
RL  Nat. Biotechnol. 21:1307-1313(2003).
DR  EMBL; BX571862; CAE13380.1; -.
DR  Photolyst; plul085; -.
KW  Complete proteome.
SQ  SEQUENCE 960 AA; 107431 MW; 1329711136B972B1 CRC64;

Query Match 67.3%; Score 37; DB 2; Length 960;
Best Local Similarity 77.8%; Pred. No. 1.6e+02;
Matches 7; Conservative 1; Mismatches 1; Indels 0; Gaps 0;

Qy  2 RLTRKDRGL 10
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Db  646 RLTRADRCM 654

RESULT 15
URE3 CORGL STANDARD; PRT; 100 AA.
AC  Q9RHM6;
DT  29-MAR-2004 (Rel. 43, Last sequence update)
DT  29-MAR-2004 (Rel. 43, Last sequence update)
DT  05-JUL-2004 (Rel. 44, Last annotation update)
DE  Urease gamma subunit (EC 3.5.1.5) (Urea amidohydrolase gamma subunit).
GN  Name=ureA; OrderedLocustNames=Cgl0084, cgl0113;
OS  Corynebacterium glutamicum (Brevibacterium flavum).
OC  Bacteria; Actinobacteria; Actinobacteridae; Actinomycetales;
OC  Corynebacterineae; Corynebacteriaceae; Corynebacterium.
OX  NCBI_TaxID=1718;
RN  [1]
RP  SEQUENCE FROM N.A.
RC  STRAIN=ATCC 13869;
RA  Pubkas L.G., Inui M., Yukawa H.;
RT  "Structure and transcriptional regulation of the urease operon of
RT  Corynebacterium glutamicum.";
RL  Submitted (JUN-1999) to the EMBL/GenBank/DBJ databases.
RN  [2]
RP  SEQUENCE FROM N.A.
RC  STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA  Nolden L., Beckers G., Moeckel B., Nampoothiri M., Pfeifferle W.,
RA  Kraemer R., Burkovski A.;
RT  "Urease of Corynebacterium glutamicum: sequence and organisation of
RT  corresponding genes and investigation of activity.";
RL  Submitted (DEC-1999) to the EMBL/GenBank/DBJ databases.
RN  [3]
RP  SEQUENCE FROM N.A.
RC  STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RA  Nakagawa S.;
RT  "Complete genomic sequence of Corynebacterium glutamicum ATCC 13032.";
RL  Submitted (MAY-2002) to the EMBL/GenBank/DBJ databases.
RN  [4]
RP  SEQUENCE FROM N.A.
RC  STRAIN=ATCC 13032 / DSM 20300 / NCIB 10025;
RX  PubMed=12948626; DOI=10.1016/S0168-1656(03)00154-8;
RA  Kalinowski J., Bathe B., Bartels D., Bischoff N., Bott M.,
RA  Burkovski A., Busch N., Eggeing L., Eikmanns B.J., Gaigalat L.,
RA  Goesmann A., Hartmann M., Huthmacher K., Kraemer R., Linke B.,
RA  McHardy A.C., Meyer F., Moeckel B., Pfeifferle W., Puhler A.,
RA  Rey D.A., Rueckert C., Rupp O., Sahm H., Wendisch V.F., Wiegand I.,
RA  Tauch A.;
RT  "The complete Corynebacterium glutamicum ATCC 13032 genome sequence
RT  and its impact on the production of L-aspartate-derived amino acids
RT  and vitamins.";
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J. Biotechnol. 104:5-25(2003).
-|- CATALYTIC ACTIVITY: Urea + H(2)O = CO(2) + 2 NH(3).
-|- SUBUNIT: (Alpha, beta, gamma)(3) (By similarity).
-|- SUBCELLULAR LOCATION: Cytoplasmic (By similarity).
-|- SIMILARITY: Belongs to the urease gamma subunit family.

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or send an email to license@isb-sib.ch).

DR  EMBL; AB029154; BAA88552.1; -.
DR  EMBL; AJ251883; CAB81935.1; -.
DR  EMBL; AP005274; BAB97477.1; -.
DR  EMBL; BX927148; CAF18652.1; -.
DR  HSSP; P41022; 1UBP.
DR  HAMAP; MF_00739; -.
DR  InterPro; IPR002026; Urease_gamma.
DR  Pfam; PF00547; Urease_gamma_1.
DR  ProDom; PD002319; Urease_gamma_1.
DR  TIGRFAMs; TIGR00193; urease_gam; 1.
KW  Complete proteome; Hydrolase.
SQ  SEQUENCE 100 AA; 11245 MW; A48F4DC0EABD9567 CRC64;

Query Match 65.5%; Score 36; DB 1; Length 100;
Best Local Similarity 100.0%; Pred. No. 18;
Matches 7; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy  5 RKDRGLK 11
    |||||
Db  23 RKDRGLK 29

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